

**PCT**WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International BureauUSAN 11/501,464  
For IDS filed 11/10/06

## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<b>(51) International Patent Classification <sup>5</sup>:</b> <b>A61K 31/42, C07D 263/56</b>	<b>A1</b>	<b>(11) International Publication Number:</b> <b>WO 94/14435</b> <b>(43) International Publication Date:</b> 7 July 1994 (07.07.94)
<b>(21) International Application Number:</b> PCT/US93/12638 <b>(22) International Filing Date:</b> 29 December 1993 (29.12.93) <b>(30) Priority Data:</b> 07/997,818 29 December 1992 (29.12.92) US <b>(60) Parent Application or Grant</b> <b>(63) Related by Continuation</b> US 07/997,818 (CIP) Filed on 29 December 1992 (29.12.92) <b>(71) Applicant (for all designated States except US):</b> RHONE-POULENC RORER PHARMACEUTICALS INC. [US/US]; 500 Arcola Road, P.O. Box 1200, Collegeville, PA 19426-0107 (US). <b>(72) Inventors; and</b> <b>(75) Inventors/Applicants (for US only):</b> ULLRICH, John, W. [US/US]; 62 Aspen Way, Schwenksville, PA 19473 (US). KIESOW, Terence, J. [US/US]; 132 First Avenue, Pottstown, PA 19464 (US). NEUENSCHWANDER, Kent, W. [US/US]; 103 Creekside Way, Schwenksville, PA 19473 (US). SCOTESE, Anthony, C. [US/US]; 240 Beidler Road, King of Prussia, PA 19406 (US). LEARN, Keith, S.		<b>[US/US];</b> 3039 Gottschall Road, Perkiomenville, PA 18074 (US). DANKULICH, William, P. [US/US]; 230 Freeland Drive, Collegeville, PA 19426 (US). <b>(74) Agents:</b> NICHOLSON, James, A. et al.; Rhone-Poulenc Rorer, Inc., 500 Arcola Road, P.O. Box 1200, Collegeville, PA 19426-0107 (US). <b>(81) Designated States:</b> AT, AU, BB, BG, BR, CA, CH, CZ, DE, DK, ES, FI, GB, HU, JP, KP, KR, LK, LU, MG, MN, MW, NL, NO, NZ, PL, PT, RO, RU, SD, SE, SK, UA, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).  <b>Published</b> <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
<b>(54) Title:</b> CYCLOALKYL AMINE BIS-ARYL SQUALENE SYNTHASE INHIBITORS		
<b>(57) Abstract</b> <p>This invention relates to a class of novel polycyclic compounds containing a cycloalkyl ring having substituted thereon a primary amine and which is further linked or bridged to two mono- and/or bicyclic rings and which reduces levels of serum cholesterol in the body without significantly reducing mevalonic metabolite synthesis. This invention relates also to pharmacological compositions and method of treatment for lowering serum cholesterol levels using the compounds of this invention.</p> <p><b>CHOLESTEROL BIOSYNTHETIC PATHWAY</b></p>		

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	GB	United Kingdom	MR	Mauritania
AU	Australia	GE	Georgia	MW	Malawi
BB	Barbados	GN	Guinea	NE	Niger
BE	Belgium	GR	Greece	NL	Netherlands
BF	Burkina Faso	HU	Hungary	NO	Norway
BG	Bulgaria	IE	Ireland	NZ	New Zealand
BJ	Benin	IT	Italy	PL	Poland
BR	Brazil	JP	Japan	PT	Portugal
BY	Belarus	KE	Kenya	RO	Romania
CA	Canada	KG	Kyrgyzstan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic of Korea	SD	Sudan
CG	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SI	Slovenia
CI	Côte d'Ivoire	LI	Liechtenstein	SK	Slovakia
CM	Cameroon	LK	Sri Lanka	SN	Senegal
CN	China	LU	Luxembourg	TD	Chad
CS	Czechoslovakia	LV	Latvia	TG	Togo
CZ	Czech Republic	MC	Monaco	TJ	Tajikistan
DE	Germany	MD	Republic of Moldova	TT	Trinidad and Tobago
DK	Denmark	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	US	United States of America
FI	Finland	MN	Mongolia	UZ	Uzbekistan
FR	France			VN	Viet Nam
GA	Gabon				

5

## CYCLOALKYL AMINE BIS-ARYL SQUALENE SYNTHASE INHIBITORS

10

Field of the Invention

The present invention relates to a class of novel compounds useful in the treatment of diseases associated with undesirable cholesterol levels in the body, and particularly diseases of the cardiovascular system, such as atherosclerosis.

15

Only about 7% of the total body cholesterol circulates in the plasma, where it has been linked to atherosclerosis. The remaining 93% is located in cells, where it performs vital structural and metabolic functions. Excluding the diet, which accounts for approximately one-third of the total body cholesterol, the cells obtain the necessary cholesterol by endogenous biosynthesis (Figure 1) or by removing low density lipoprotein (LDL) from the bloodstream. Approaches to the control of plasma cholesterol levels have been varied, however it has been shown that inhibiting endogenous cholesterol biosynthesis forces the cell to rely more on LDL uptake to satisfy their cholesterol requirements. Increased LDL uptake by cells, especially liver cells, has been shown to lower plasma cholesterol levels.

Squalene synthase is a microsomal enzyme that catalyzes the reductive dimerization of two molecules of farnesyl diphosphate to form squalene. While farnesyl diphosphate serves as the precursor to several other biologically important compounds, squalene is utilized only for cholesterol biosynthesis. Consequently, this is the first totally committed step in the biosynthesis of cholesterol (see Figure 1). Inhibition at this step would stop only *de novo* cholesterol synthesis while allowing other essential pathways to isopentenyl tRNA, the prenylated proteins, ubiquinone, and dolichol to proceed unimpeded.

Inhibition of HMG-CoA reductase, an enzyme positioned early in the cholesterol biosynthetic pathway, results in a decrease of *de novo* cholesterol biosynthesis and an accompanying up-regulation of LDL receptors. However due to a large induction in the amount of the HMG-CoA reductase enzyme, the effect of this inhibition is blunted somewhat and the maximum LDL cholesterol reductions attainable are limited. Since inhibition of squalene synthase does not cause the same amount of enzyme induction (HMG-CoA reductase or squalene synthase), it causes in a greater reduction of *de novo* cholesterol biosynthesis. This translates into more up-regulation of LDL receptors than is seen with an HMG-CoA reductase inhibitor and greater efficacy for lowering circulating LDL levels.

#### Reported Developments

The literature describes the cholesterol biosynthetic pathway and possible means for the inhibition of squalene synthase. In a series of papers including J. Am. Chem. Soc., 1982, 104, 7376-7378 and J. Am. Chem. Soc., 1989, 111, 3734-3739, C. Dale Poulter, et al disclose that ammonium substituted cyclopropyl polyene compounds mimic the topological and electrostatic properties of the primary cation and tertiary cation of presqualene diphosphate. and in the presence of phosphate buffer, inhibit squalene synthase. Scott A. Biller et al in J. Med. Chem., 1988, 31, 1869-1871 disclose that a series of stable, non-ionizable analogues of farnesyl diphosphate, comprising phosphomethylene phosphate polyene compounds, inhibit squalene synthase.

International Patent Application published under the Patent Cooperation Treaty having International Publication Number: WO 92/15579 and assigned to the same assignee as the present application, is directed to multicyclic tertiary amine polyaromatic squalene synthetase inhibitors. These compounds all contain a multicyclic ring having a nitrogen atom therein.

United States Patent 5,135,935 assigned to Merck and Co., is directed to squalene synthetase inhibitors which are aryl-oxadiazole-quinuclidines. International Patent Applications published under the Patent Cooperation Treaty having International Publication Numbers: WO 92/12159, 92/12158, 92/12157, 92/12156 and 92/12160 and being assigned to Glaxo Group Ltd. are directed to

bridged cyclic ketal derivatives for lowering the level of blood plasma cholesterol.

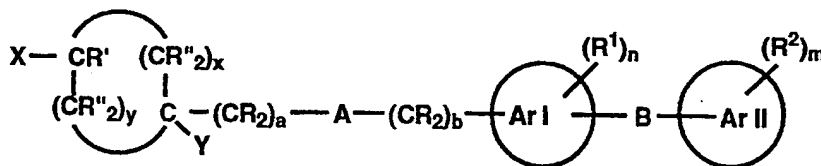
5       The present invention is directed to a class of novel cycloalkyl-diaryl compounds having a primary amine substituted on the cycloalkyl ring and which exhibit squalene synthase inhibition properties.

## Summary of the Invention

10 This invention comprises polycyclic compounds containing a cycloalkyl ring having substituted thereon a primary amine and which is further linked or bridged to two mono- and/or bicyclic rings. The compounds of this invention possess properties which reduce levels of serum cholesterol in the body without significantly reducing mevalonic metabolite synthesis and thus provide a  
15 therapeutic agent having fewer side effects than agents which act by inhibiting the HMG-CoA reductase enzyme. This invention relates also to pharmacological compositions and method of treatment for lowering serum cholesterol levels using the compounds of this invention.

20        The compounds of the present invention are described as amino cycloalkyl-bis aryl ring compounds containing two linked or bridged mono- and/or bicyclic aryl rings and are further linked or bridged to a cycloalkyl ring having a primary amine attached thereon.

25 More specifically, the compounds of this invention are described by the compounds of Formula I:



### Formula I

30                    where:

Ar I and Ar II are independently a mono- or di- aryl or heteroaryl ring;

A is O, S, NR, SO, SO<sub>2</sub>, NR-C=O, O=C—NR, RC=CR, C<sub>j</sub>/C or a bond;

B is  $(CR_2)_e$ , O, S, NR, SO,  $SO_2$ ,  $NR-C=O$ ,  $O=C-NR$ ,  $RC=CR$ ,  $C\equiv C$ ,  $O=C$  or a bond when Ar II is mono- aryl or heteroaryl and  $(CR_2)_e$ , O, S,  $O=C$ , NR, SO,  $SO_2$  or a bond when Ar II is di- aryl or heteroaryl;

5

one of X or Y is  $R''$ ;

one of X or Y is 
$$\begin{array}{c} R_\alpha \\ | \\ N - (CR_2)_z - \\ | \\ R_\beta \end{array}$$
 ;

10

a and b are independently 0-4 and  $a + b = 0-4$ ;

e is 1-2;

m and n are independently 0-2;

15

x is 1-6;

y is 0-2;

20

$x + y$  is 1-6;

z is 0-3;

$R_\alpha$  is hydrogen or alkyl;

25

$R_\beta$  is hydrogen, alkyl or hydroxy;

$R'$  is hydrogen, alkyl, hydroxy, alkoxy or halo;

30

$R''$  is hydrogen, alkyl, hydroxy, hydroxyalkyl, alkoxy, alkoxyalkyl or halo;

$R'''$  is hydrogen, alkyl, hydroxy or carboxy;

one set of vicinal  $R''$  and  $R'''$  groups may form a double bond  
35 when  $x+y=3-6$ ;

R is hydrogen or alkyl;

R<sup>1</sup> and R<sup>2</sup> are independently hydrogen, alkyl, alkoxy, hydroxy, halo,  
5 haloalkyl, nitro, amino or mono- or di-alkylamino; and

its stereoisomers, enantiomers, diastereoisomers and racemic mixtures; or  
pharmaceutically acceptable salts thereof.

10

#### Description of the Drawing

Figure 1 is a schematic diagram of the biosynthetic pathway of  
cholesterol.

15

#### Detailed Description and Preferred Embodiments

As employed above and throughout this disclosure, the following terms,  
20 unless otherwise indicated, shall be understood to have the following meanings:

"Biological pH" refers to that pH of blood, plasma or serum in the body  
between about 7.2 and about 7.5 and which does not interfere with normal  
degradation of materials present therein. The normal pH of blood, plasma or  
25 serum values is about 7.35-7.45 and is preferably about pH 7.39-7.41.

"Monocyclic aryl" means a carbocyclic and/or heterocyclic aromatic ring.  
Preferred groups include phenyl, thienyl, pyridinyl, furyl and pyrimidinyl.

30 "Bicyclic aryl" means a bicyclic ring system composed of two fused  
carbocyclic and/or heterocyclic aromatic rings. Preferred groups include  
naphthyl, indolyl, benzothienyl, benzofuranyl, quinoliny, benzoxazole and  
benzothiazole.

35

"Aryl" means a carbocyclic or heterocyclic aromatic ring.

"Alkyl" means a saturated aliphatic hydrocarbon, either branched- or straight-chained. Preferred alkyl is "loweralkyl" having about 1 to about 6 carbon atoms. Examples of alkyl include methyl, ethyl, n-propyl, isopropyl, butyl, sec-butyl, t-butyl, amyl and hexyl.

5

"Alkoxy" refers to an alkyl-O-group.

"Alkenyl" refers to a hydrocarbon having at least one point of unsaturation and may be branched- or straight-chained. Preferred alkenyl groups have 2 to about 6 carbon atoms. Exemplary alkenyl groups include vinyl, allyl, ethynyl and isopropenyl.

10

The preferred aryloxy group is phenoxy.

15

"Aralkyl" means an alkyl group substituted by an aryl radical. The preferred aralkyl groups are benzyl or phenethyl.

The preferred aryloxy group is phenoxy.

20

The preferred aralkoxy groups are benzyloxy and phenethoxy.

The preferred acyloxy group is acetoxy.

"Halo" means a halogen. Preferred halogens include chloride, bromide and fluoride. The preferred haloalkyl group is trifluoromethyl.

25

Preferred compounds of Formula I are defined where Ar I and Ar II are independently a substituted or unsubstituted mono- or bicyclic system of about 5 to about 14 atoms which may be partially or completely unsaturated carbocyclic or heterocyclic and where each ring of said system contains 0 to about 2 hetero atoms selected from N, O and S provided said hetero atoms are not vicinal oxygen and/or sulfur atoms and where the substituents may be located at any appropriate position of the ring system and are described by the R definition above.

30

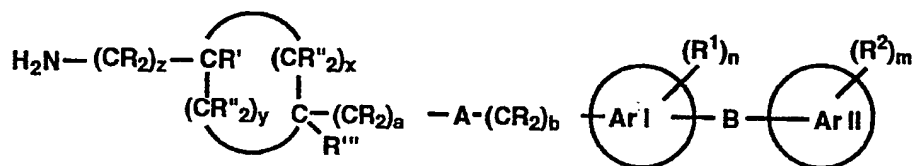
Preferred monocyclic rings include aryl and unsaturated carbocyclic and heterocyclic rings. Exemplary rings are substituted or unsubstituted pyrrole,



thiophene, furan, cyclopentadiene, imidazole, pyrazole, 1,2,4-triazole, pyridine, pyrazine, pyrimidine, pyridazine, thiazole, isothiazole, oxazole, isoxazole, s-triazine and benzene.

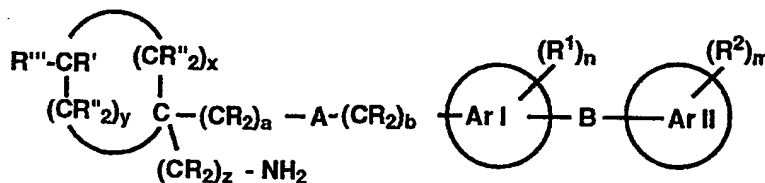
- 5 Preferred bicyclic ring systems include bicyclic aryl and bicyclic unsaturated carbocyclic and heterocyclic rings. Exemplary bicyclic rings include substituted and unsubstituted indene, isoindene, benzofuran, dihydrobenzofuran, benzothiophene, indole, 1H-indazole, indoline, imadazole; azulene, tetrahydroazulene, benzofuran, benzothiaphene, benzopyrazole, 10 benzoimidazole, benzoxazole, benzothiazole, 1,3-benzodioxole, 1,4-benzodioxan, purine, naphthalene, tetralin, coumarin, chromone, chromene, 1,2-dihydrobenzothiopyran, tetrahydrobenzothiopyran, quinoline, isoquinoline, quinazoline, pyrido[3,4-b]-pyridine, and 1,4-benzisoxazine.

- 15 The more preferred compounds of this invention include compounds of Formulae II and III:



Formula II

20

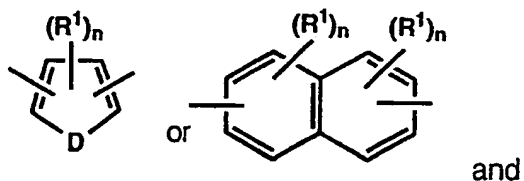


Formula III

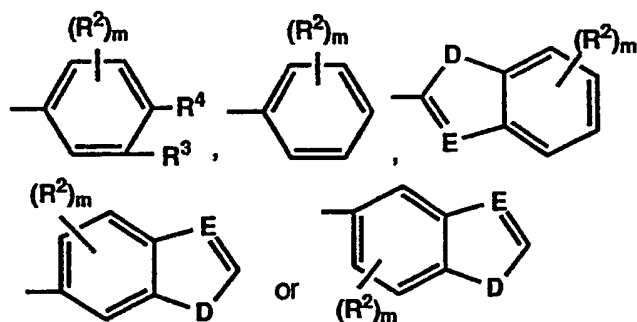
where:

25

Ar I is



Ar II is



where:

5 D is CH<sub>2</sub>, CH=CH, O, S or N-R;

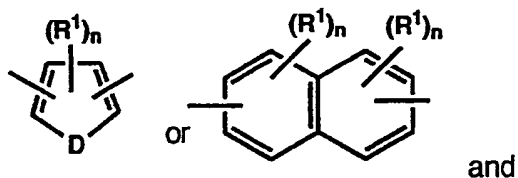
E is CH, N or CH-O;

10 R<sup>1</sup> and R<sup>2</sup> are independently hydrogen, alkyl, alkoxy, hydroxy, halo or haloalkyl;

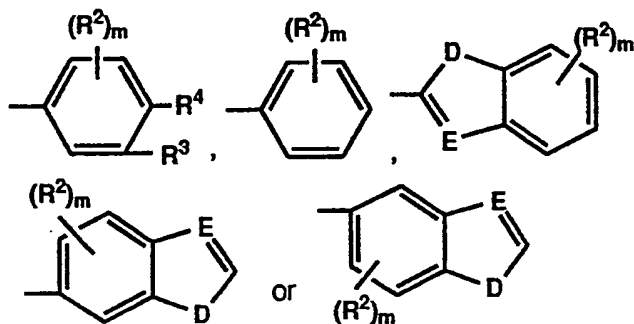
R<sup>3</sup> and R<sup>4</sup> together are O-CH<sub>2</sub>-CH<sub>2</sub>, O-CH<sub>2</sub>-CH<sub>2</sub>-CH<sub>2</sub>, O-CH<sub>2</sub>-O or O-CH<sub>2</sub>-CH<sub>2</sub>-O.

15 The more preferred compounds are those described by Formulae II and III where:

Ar I is



20 Ar II is



where:

D is CH<sub>2</sub>, CH=CH, O, S or N-R;

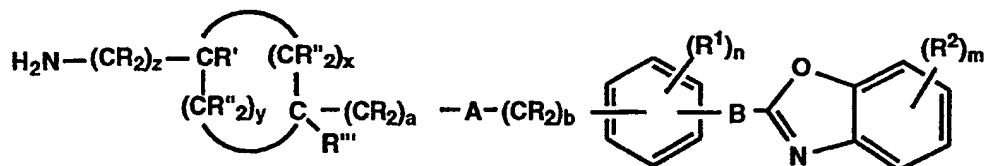
5 E is CH, N or CH-O;

R<sup>1</sup> and R<sup>2</sup> are independently hydrogen, alkyl, alkoxy, hydroxy, halo or trifluoromethyl; and

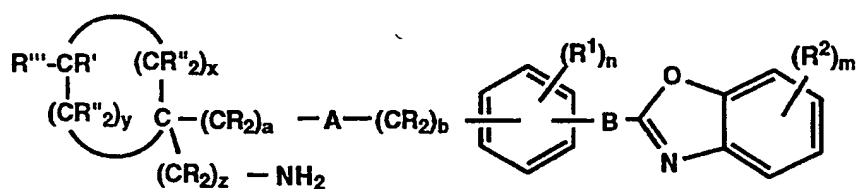
10 R<sup>3</sup> and R<sup>4</sup> together are O-CH<sub>2</sub>-CH<sub>2</sub>, O-CH<sub>2</sub>-CH<sub>2</sub>-CH<sub>2</sub>, O-CH<sub>2</sub>-O or O-CH<sub>2</sub>-CH<sub>2</sub>-O.

A special embodiment of this invention describes those compounds of Formula IV through VII:

15

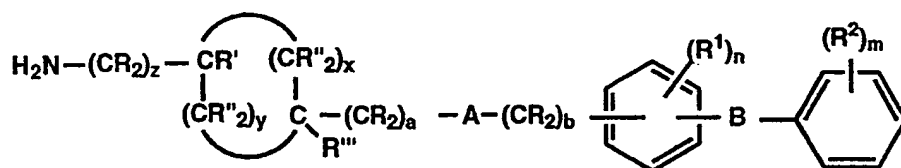


Formula IV

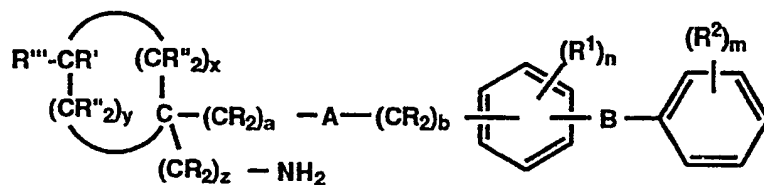


20

Formula V



Formula VI



Formula VII

where:

5           A is O, S, NR, CH=CH or a bond;

B is (CR<sub>2</sub>)<sub>e</sub>, O, S, NR, O=C, CH=CH or a bond when Ar II is mono-aryl and (CR<sub>2</sub>)<sub>e</sub>, O, S, O=C, or NR when Ar II is di- aryl;

10           a and b are independently 0-3 and a + b = 0-3;

e is 1-2;

m and n are independently 0-2;

15

x is 1-6;

y is 0-2;

20

x + y is 1-6;

z is 0-3;

R' is hydrogen, alkyl or hydroxy;

25

R'' is hydrogen, alkyl or hydroxy;

R''' is hydrogen, alkyl, hydroxy or carboxy;

30

one set of vicinal R'' and R''' groups may form a double bond when x+y=3-6;

R is hydrogen or alkyl; and

$R^1$  and  $R^2$  are independently hydrogen, alkyl, alkoxy, hydroxy, halo or trifluoromethyl.

5        The most preferred compounds described by Formulae IV-VII include those where:

A is O or a bond;

10        B is  $\text{CH}=\text{CH}$  or a bond when Ar II is mono- aryl and a bond when Ar II is di- aryl;

a is 0-1;

15        b is 0-1;

m and n are independently 0-1;

x is 1-6;

20

y is 0-2;

x + y is 1-6;

25

z is 0-1;

R,  $R'$ ,  $R''$ ,  $R'''$  and  $R^1$  are hydrogen or hydroxy; and

$R^2$  is hydrogen, alkyl, alkoxy or halo.

30

A special embodiment includes those compounds described by Formulae IV-V where:

A is O;

35        B is  $\text{CH}=\text{CH}$  or a bond when Ar II is mono- aryl and a bond when Ar II is di- aryl;

a is 0-1;

b is 0-1;

5 m and n are independently 0-1;

x is 1-6;

y is 0-2;

10

x + y is 1-6;

z is 0-1;

15

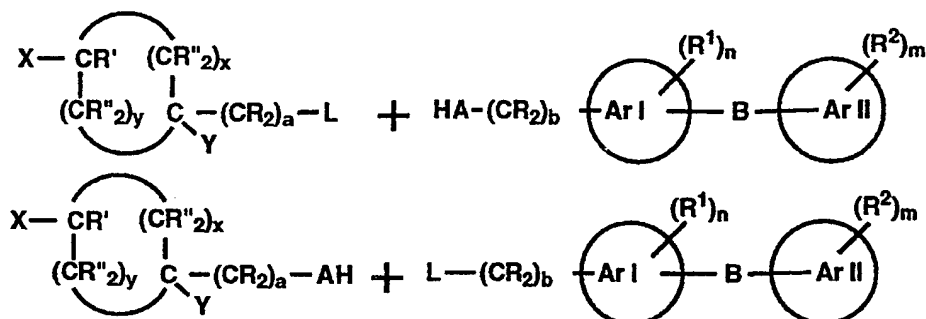
R, R', R'', R''' and R<sup>1</sup> are hydrogen or hydroxy; and

R<sup>2</sup> is hydrogen, alkyl, alkoxy or halo.

20 The compounds of this invention may be prepared by employing procedures known in the literature starting from known compounds or readily preparable intermediates Exemplary general procedures are described below.

25 It is convenient to synthesize these molecules by employing condensation reactions at the above-described reactive A and B sites of the molecule. Exemplary general procedures are shown below and, for convenience, describe those compounds where Ar I is a benzene ring and Ar II is a benzoxazole ring. Of course, while the following reactions involved are basic to developing substituted phenyl-benzoxazole molecules having desirable substituent groups present, the substitution pattern for other mono- or bicyclic rings depends on the chemistry of the particular ring. Any such adjustments to  
30 the chemistry would be familiar to one skilled in the art.

Thus, in order to prepare those compounds where A is O, S or NR and B (CR<sub>2</sub>)<sub>e</sub>, O, S, NR, SO, SO<sub>2</sub>, NR—C=O, O=C—NR, RC=CR, C≡C, O=C or  
35 a bond, the following reactions or combination of reactions are employed:



where L is a leaving group, preferably halo, tosylate or mesylate.

The amine is protected with the usual protecting groups such as  
 5 phthalimide, triphenylmethyl or hydroborane complex which is removed at the appropriate time.

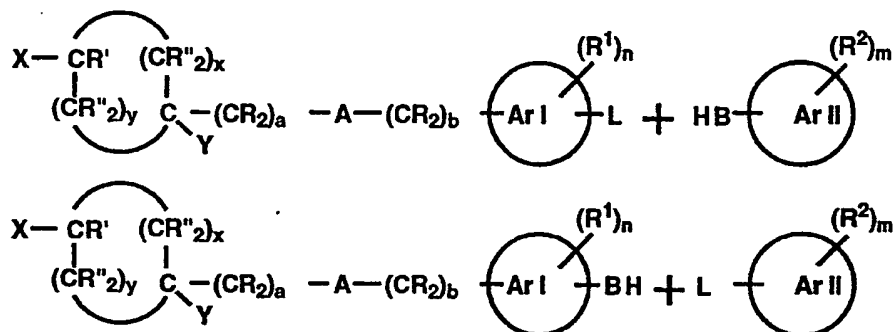
Where A is O or S, any base normally employed to deprotonate an  
 alcohol or thiol may be used, such as sodium hydroxide, potassium hydroxide,  
 10 sodium hydride, triethyl amine, sodium bicarbonate or diisopropylethylamine.

Reaction temperatures are in the range of -78°C to reflux depending on  
 the reactants involved. (Preferably about 0°C to room temperature). Reaction  
 times vary from about 2 to about 96 hours. The reaction is usually carried out in  
 15 a solvent that will dissolve both reactants and is inert to both as well. Solvents  
 include, but are not limited to, diethyl ether, tetrahydrofuran, N,N-dimethyl  
 formamide, dimethyl sulfoxide, dioxane and the like.

In the case where A or B is SO or SO<sub>2</sub>, then treatment of the thio  
 20 compound with m-chlorobenzoic acid or sodium periodate results in the sulfinyl  
 compound. Preparation of the sulfonyl compound may be accomplished by  
 known procedures such as dissolving the sulfinyl compound in acetic acid and  
 treating hydrogen peroxide, preferably about 30% aqueous H<sub>2</sub>O<sub>2</sub>.

In certain of the reaction schemes a metal salt may be used. Any  
 25 appropriate metal salt formed from such as Li, K, Na, Mg, Br or the like may be  
 used.

Those compounds where A is (CR<sub>2</sub>)<sub>e</sub>, RC=CR, O, S, C≡C, or a bond,  
 30 may be prepared by one of the following reactions:



where L is a leaving group, preferably halo, tosylate or mesylate.

- 5           The amine of course, is protected by a suitable protecting group such as phthalimide, triphenylmethyl or hydroborane complex which is removed at the appropriate time. These reaction steps to protect the various substituents involved will be by known methods in the art. Further, reaction conditions may be employed as determined by the particular Ar I and Ar II rings involved as well as the presence of any reactive R'', R''', R<sup>1</sup> or R<sup>2</sup> group.
- 10

Various substituents on the present new compounds can be present in the starting compounds, added to any one of the intermediates or added after formation of the final products by known methods of substitution or conversion reactions. If the substituents themselves are reactive, then the substituents can themselves be protected according to the techniques known in the art. A variety of protecting groups known in the art, may be employed. Examples of many of these possible groups may be found in "Protective Groups in Organic Synthesis" by T.W. Green, John Wiley and Sons, 1981. For example, nitro groups can be added by nitration and the nitro group converted to other groups, such as amino by reduction, and halo by diazotization of the amino group and replacement of the diazo group. Acyl groups can be added by Friedel-Crafts acylation. The acyl groups can then be transformed to the corresponding alkyl groups by various methods, including the Wolff-Kishner reduction and Clemmenson reduction. Amino groups can be alkylated to form mono- and di-alkylamino groups; and mercapto and hydroxy groups can be alkylated to form corresponding ethers. Primary alcohols can be oxidized by oxidizing agents known in the art to form carboxylic acids or aldehydes, and secondary alcohols can be oxidized to form ketones. Thus, substitution or alteration reactions can

15

20

25



be employed to provide a variety of substituents throughout the molecule of the starting material, intermediates, or the final product.

Since the compounds of this invention have certain substituents which  
5 are necessarily present, the introduction of each substituent is, of course,  
dependent on the specific substituents involved and the chemistry necessary for  
their formation. Thus, consideration of how one substituent would be affected  
by a chemical reaction when forming a second substituent would involve  
techniques familiar to the skilled artisan. This would further be dependent on  
10 the ring involved.

Certain compounds of this invention may have at least one asymmetric  
carbon atom such as those compounds having different geminal groups or those  
compounds which contain an asymmetric carbon atom. Further, certain  
15 compounds of this invention may exist in their cis or trans configuration such as  
those compounds where A or B is CR=CR or R<sup>n</sup> and R<sup>m</sup> form a double bond. As  
a result, those compounds of this invention may be obtained either as racemic  
mixtures, diastereoisomeric mixtures or as individual enantiomers. When two or  
three asymmetric centers are present the product may exist as mixtures of two  
20 or four diastereomers. Of course it is understood that certain other compounds  
within the scope of this invention could have a number of stereocenters. In  
general, a compound with x stereocenters can have a maximum of 2<sup>x</sup>  
stereoisomers. Therefore, a compound having three such centers gives rise to  
a maximum of eight stereoisomers, while one having four produces sixteen, etc.  
25 The product may be synthesized as a mixture of the isomers and then the  
desired isomer separated by conventional techniques such as chromatography  
or fractional crystallization from which each diastereomer may be resolved. On  
the other hand, synthesis may be carried out by known stereospecific processes  
using the desired form of the intermediate which would result in obtaining the  
30 desired stereospecificity.

Reference to the separation of cis and trans isomers by chromatography  
may be found in W.K. Chan, et al, J. Am. Chem. Soc. 96, 3642, 1974.

35 It is to be understood that the scope of this invention encompasses not  
only the various isomers which may exist but also the various mixture of isomers  
which may be formed.

The resolution of the compounds of this invention and their starting materials may be carried out by known procedures. Incorporation by reference is hereby made to the four volume compendium Optical Resolution Procedures for Chemical Compounds: Optical Resolution Information Center, Manhattan College, Riverdale, New York. Such procedures are useful in the practice of this invention. A further useful reference is Enantiomers, Racemates and Resolutions: Jean Jacques, Andre Collet and Samuel H. Wilen; John Wiley & Sons, Inc., New York, 1981. Basically, the resolution of the compounds is based on the differences in the physical properties of diastereomers. Conversion of the racemates into a mixture of diastereomers by attachment of an enantiomerically pure moiety results in forms that are separable by fractional crystallization, distillation or chromatography.

The present compounds form salts with acids when a basic amino function is present and salts with bases when an acid function, i.e., carboxyl, is present. All such salts are useful in the isolation and/or purification of the new products. Of particular value are the pharmaceutically acceptable salts with both acids and bases. Suitable acids include, for example, hydrochloric, sulfuric, nitric, benzenesulfonic, toluenesulfonic, acetic, maleic, tartaric and the like which are pharmaceutically acceptable. Basic salts for pharmaceutical use are the Na, K, Ca and Mg salts.

The compounds of the present invention may be prepared by the following representative examples.

#### Example 1

##### Trans-2-[4-(benzoxazol-2-yl)benzyloxy]cyclohexylamine acetate

#### **Step A:** Trans-2-(N-phthaloyl)cyclohexanol

A mixture of trans-2-aminocyclohexanol (50 g, 0.33 mol), phthalic anhydride (58 g, 0.39 mol) and triethylamine (160.8 ml, 1.15 mol) in 500 ml chloroform is stirred at reflux under nitrogen overnight. The solution is then cooled to room temperature, washed with aqueous sodium carbonate, the chloroform layer is separated, dried and concentrated in vacuo. The crude

compound is passed through silica gel using ethylene chloride as eluent to obtain trans-2-(N-phthaloyl)-cyclohexanol which is used directly in the next step.

**Step B:**     Trans-1-[4-(benzoxazol-2-yl)benzyloxy]-2-phthaloylcyclohexane

5

To an activated mixture of sodium hydride (60% in mineral oil; 0.72 g, 18 mmol) in 20 ml DMF is added trans-2-(N-phthaloyl)cyclohexanol (4 g; 16.32 mmol) in 20 ml DMF, dropwise. The solution is stirred under nitrogen at room temperature for 1 hr. To this is added dropwise 4-(benzoxazol-2-yl)benzyl bromide (5.2 g; 18.00 mmol) in 16 ml DMF under nitrogen. The solution is stirred at room temperature overnight. The solution is then poured into ice/ammonium chloride/ethyl-acetate/hexane. The organic layer is separated, washed with H<sub>2</sub>O: brine ; 1:1, 5X, dried, concentrated in vacuo and passed through a silica gel column using 0-2% MeOH:CH<sub>2</sub>Cl<sub>2</sub> to obtain trans-1-[4-(benzoxazol-2-yl)benzyloxy]-2-phthaloyl-cyclohexane which is used directly in the next step.

10

15

**Step C:**     Trans-2-[4-(benzoxazol-2-yl)benzyloxy]cyclohexylamine acetate

20

To a mixture of trans-1-[4-(benzoxazol-2-yl)benzyloxy]-2-phthaloylcyclohexane (3.7 g; 8.18 mmol), in isopropanol (70 ml) and water (13 ml) is added sodium borohydride (1.5 g; 40.93 mmol). The solution is stirred at room temperature under nitrogen overnight. Acetic acid is then slowly added to the reaction mixture at room temperature and then the temperature of the mixture is allowed to rise to 110°C. The reaction mixture is stirred with refluxing under nitrogen overnight. The solution is then cooled to room temperature and azeotroped 6x with toluene and the evacuated. The crude material is purified over silica gel with 0-20% MeOH:CH<sub>2</sub>Cl<sub>2</sub> and recrystallized from EtOH/hexane to obtain trans-2-[4-(benzoxazol-2-yl)benzyloxy]cyclohexylamine acetate.

25

30

## Example 2

Cis-2-[(4-benzoxazol-2-yl)benzyloxy]cyclohexylamine hydrochloride

35

**Step A:**     Cis-2-triphenylmethylaminocyclohexanol

A mixture of cis-2-aminocyclohexanol hydrochloride (4.88 g; 32 mmol) (William S. Johnson and Elliot N. Schubert, J.A.C.S., Vol. 72, pp. 2187-2190 (1950), triphenylmethylchloride (10.7 g; 38.4 mmol) and triethylamine (16 ml; 115 mmol) in 250 ml methylene chloride are refluxed for 3 hrs. then stirred at  
5 room temperature overnight. The mixture is poured into H<sub>2</sub>O and extracted with CH<sub>2</sub>Cl<sub>2</sub> (200 ml), dried (Na<sub>2</sub>SO<sub>4</sub>) concentrated in vacuo and flash chromatographed using 9:1  $\Rightarrow$  7:1; hexane:EtOAc to obtain cis-2-triphenylmethylaminocyclohexanol which is used directly in the next step.

10 **Step B:** Cis-1-[4-(benzoxazol-2-yl)benzyloxy]-2-triphenylmethylamino-  
cyclohexane

Cis-2-triphenylmethylaminocyclohexanol (1.52 g; 4.25 mmol) is added to a solution of sodium hydride (256 mg; 60%; 6.4 mmol) in 20 ml dry THF and  
15 refluxed overnight. The solution is cooled, 4-(benzoxazol-2-yl)benzyl bromide (1.3g; 4.5 mmol) is added in one portion and refluxed for 5 hrs. This is then poured into water and extracted with 2x100 ml CH<sub>2</sub>Cl<sub>2</sub>, dried (Na<sub>2</sub>SO<sub>4</sub>), concentrated to dryness and flash chromatographed using 4:1  $\Rightarrow$  8:1; hexane:EtOAc to obtain cis-1-[4-(benzoxazol-2-yl)benzyloxy]-2-triphenylmethyl-  
20 aminocyclohexane which is used directly in the next step.

**Step C:** Cis-2-[(4-benzoxazol-2-yl)benzyloxy]cyclohexylamine  
hydrochloride

25 Cis-1-[4-(benzoxazol-2-yl)benzyloxy]-2-triphenylmethylaminocyclohexane (2.03 g; 3.6 mmol) in 200 ml ethanol is treated with ethanolic HCl to pH 4 and stirred for 24 hrs. at room temperature. The solution is concentrated in vacuo, treated with ether and a white solid which precipitates out is filtered, washed with dry ether and dried in vacuo to obtain cis-2-[(4-benzoxazol-2-yl)benzyloxy]-  
30 cyclohexylamine hydrochloride (m. p. 280-281°C dec).

**Example 3**

35 2 $\alpha$ -hydroxy-3 $\beta$ [(4-benzoxazol-2-yl)benzyloxy]cyclohexyl-1 $\beta$ -amine

2 $\beta$ -hydroxy-3 $\beta$ [(4-benzoxazol-2-yl)benzyloxy]cyclohexyl-1 $\alpha$ -amine

**Step A:** 3-[3-(benzoxazol-2-yl)benzyloxy]cyclohexane

5 A mixture of 2-cyclohexan-1-ol (0.63 ml; 6.4 mmol) and sodium hydride (60%; 384 g; 9.6 mmol) in 30 ml dry THF is refluxed overnight. The solution is cooled and 4-(benzoxazol-2-yl)benzyl bromide (1.95 g; 6.75 mmol) is added in one portion and refluxing continued for 5 hrs. The solution is quenched with the addition of 2 ml H<sub>2</sub>O, concentrated in vacuo poured into H<sub>2</sub>O and extracted with 2 x 100 ml CH<sub>2</sub>Cl<sub>2</sub>, dried (Na<sub>2</sub>SO<sub>4</sub>), concentrated in vacuo and flash  
10 chromatographed using 9:1  $\Rightarrow$  6:1 ; hexane:EtOAc to obtain 3-[3-(benzoxazol-2-yl)benzyloxy]cyclohexane which is used directly in the next step.

**Step B:** 3-(benzoxazol-2-yl)benzyloxycyclohex-1 $\beta$ -an-2 $\alpha$ ,3 $\alpha$ -epoxide

15 3-(benzoxazol-2-yl)benzyloxycyclohex-1 $\alpha$ -an-2 $\alpha$ ,3 $\alpha$ -epoxide

3-[3-(Benzoxazol-1-yl)benzyloxy]cyclohexane (1.26 g; 4.1 mmol) is added to m-chloroperbenzoic acid (973 mg; 80%; 4.5 mmol) in dry methylene chloride (100 ml) at 0°C. The solution is allowed to warm slowly to room temperature  
20 and stirred overnight. The solution is then poured into H<sub>2</sub>O and extracted 2 x 100 ml CH<sub>2</sub>Cl<sub>2</sub>, dried (Na<sub>2</sub>SO<sub>4</sub>), concentrated in vacuo and flash chromatographed using 6:1  $\Rightarrow$  3:1 ; hexane:EtOAc to afford two products. The product having the higher RF values and weighing 0.87 g is the cis-epoxide, 3-(benzoxazol-2-yl)benzyloxy-cyclohex-1 $\alpha$ -an-2 $\alpha$ ,3 $\alpha$ -epoxide and that having the  
25 lower RF values and weighing 0.35 g is the trans-epoxide, 3-(benzoxazol-2-yl)benzyloxycyclohex-1 $\beta$ -an-2 $\alpha$ ,3 $\alpha$ -epoxide which is verified by N M R. These products are used directly in the next steps.

**Step C:** 2 $\alpha$ -hydroxy-3 $\beta$ [(4-benzoxazol-2-yl)benzyloxy]cyclohexyl-1 $\beta$ -amine

30

3-(Benzoxazol-2-yl)benzyloxycyclohex-1 $\beta$ -an-2 $\alpha$ ,3 $\alpha$ -epoxide, (330 mg) is combined with NH<sub>4</sub>OH (2 ml) in MeOH (8 ml) and heated in a sealed tube at 80°C for 12 hrs. The solution is then concentrated in vacuo and subjected to flash chromatography using a gradient elution of MeOH  $\Rightarrow$  5% Et<sub>3</sub>N/MeOH to  
35 afford 2 $\alpha$ -hydroxy-3 $\beta$ [(4-benzoxazol-2-yl)benzyloxy]cyclohexyl-1 $\beta$ -amine (m. p. 147-148°C).

**Step D:** 2 $\beta$ -hydroxy-3 $\beta$ [(4-benzoxazol-2-yl)benzyloxy]cyclohexyl-1 $\alpha$ -amine

Following the procedure of Step C, above, and replacing 3-(benzoxazol-2-yl)benzyloxycyclohex-1 $\beta$ -an-2 $\alpha$ ,3 $\alpha$ -epoxide with 3-(benzoxazol-2-yl)-  
5 benzyloxy-cyclohex-1 $\alpha$ -an-2 $\alpha$ ,3 $\alpha$ -epoxide then the product prepared is 2 $\beta$ -hydroxy-3 $\beta$ [(4-benzoxazol-2-yl)benzyloxy]cyclohexyl-1 $\alpha$ -amine.

**Example 4**

10

Trans-2-amino-[4-(benzoxazol-2-yl)benzyloxy]cyclopentane hydrochloride

**Step A:** Trans-2-aminocyclopentanol

15 A mixture of cyclopentene oxide (4 ml; 45.84 mmol), ammonium hydroxide solution (10 ml) and 4 ml methanol are heated in a sealed tube at 80°C overnight. The reaction mixture is cooled, concentrated in vacuo to dryness and used directly in the next step.

20 **Step B:** Trans-2-triphenylmethylaminocyclopentanol

A mixture of trans-2-aminocyclopentanol (45.84 mmol), triphenylmethylchloride (11.50 g; 41.26 mmol) and triethylamine (15.97 ml; 114.60 mmol) in 100 ml CH<sub>2</sub>Cl<sub>2</sub> is refluxed overnight. The reaction mixture is cooled and  
25 concentrated in vacuo and flash chromatographed using 5:1 ; hexane:EtOAc to obtain trans-2-triphenylmethylaminocyclopentanol which is used directly in the next step.

**Step C:** Trans-2-triphenylmethylamino-[4-(benzoxazol-2-yl)benzyloxy]cyclopentane  
30

To a mixture of trans-2-triphenylmethylaminocyclopentanol (4.76 g; 13.85 mmol), is added NaH (60%; 0.83 g; 20.78 mmol) in 100 ml THF and 20 ml DMF. This mixture is stirred for 1 hr. and then 4-(benzoxazol-2-yl)benzyl bromide (3.99  
35 g; 13.85 mmol) is added followed by potassium iodide (pinch). The reaction mixture is stirred for 72 hrs., quenched with water, concentrated in vacuo and flash chromatographed using 9:1 ; hexane:EtOAc to obtain trans-2-

triphenylmethylamino-[4-(benzoxazol-2-yl)benzyloxy]cyclopentane which is used directly in the next step.

5      **Step D:**      Trans-2-amino-[4-(benzoxazol-2-yl)benzyloxy]cyclopentane  
hydrochloride

To a mixture of trans-2-triphenylmethylamino-[4-(benzoxazol-2-yl)benzyloxy]-cyclopentane (3.67 g; 6.67 mmol) in ethanol (100 ml) and CH<sub>2</sub>Cl<sub>2</sub> (20 ml) is added ethanolic HCl until pH=2. This is stirred overnight,  
10 concentrated in vacuo, washed with CH<sub>2</sub>Cl<sub>2</sub>, dried under vacuum and flash chromatographed using 3:1 ; CH<sub>2</sub>Cl<sub>2</sub>:EtOH. The product is rechromatographed using 3:1 ; CH<sub>2</sub>Cl<sub>2</sub>:EtOH to obtain trans-2-amino-[4-(benzoxazol-2-yl)benzyloxy]cyclopentane hydrochloride (m.p. > 200°C).

15

**Example 5**

Trans-3-amino-[4-(benzoxazol-2-yl)benzyloxy]cyclopentane hydrochloride

20      **Step A:**      Cis-3-[benzoxazol-2-yl]benzyloxycyclopentanol

A mixture of 1,3-cyclopentanediol (5 g; 48.6 mmol), and NaH (2.15 g; 60%; 53.8 mmol) in 100 ml dry THF is refluxed overnight. This mixture is then cooled and 4-(benzoxazol-2-yl)benzyl bromide (10 g; 48.6 mmol) is added and  
25 refluxed an additional 5 hrs. To this is added 2 ml of H<sub>2</sub>O, concentrated, and then poured into H<sub>2</sub>O, extracted 2 x 100 ml CH<sub>2</sub>Cl<sub>2</sub>, dried (Na<sub>2</sub>SO<sub>4</sub>), concentrated in vacuo and flash chromatographed using CH<sub>2</sub>Cl<sub>2</sub> ⇒ 5% MeOH (CH<sub>2</sub>Cl<sub>2</sub>) to obtain a mixture of cis and trans. cis-3-[benzoxazol-2-yl]benzyloxy-cyclopentanol which is rechromatographed using HPLC with 0.25% EtOH  
30 (chloroform) as eluent to obtain cis-3-[benzoxazol-2-yl]benzyloxycyclopentanol (m.p. 182-3°C).

**Step B:**      Cis-3-chloro-[4-benzoxazol-2-yl]benzyloxy]cyclopentane

35      A mixture of cis-3-[benzoxazol-2-yl]benzyloxycyclopentanol (0.99 g; 3.2 mmol) and thionyl chloride (0.36 ml; 4.8 mmol) in dry CH<sub>2</sub>Cl<sub>2</sub> is stirred at room temperature for 72 hrs. This is then poured into water, extracted with 2x100 ml

CH<sub>2</sub>Cl<sub>2</sub>, dried (Na<sub>2</sub>SO<sub>4</sub>), concentrated in vacuo and flash chromatographed using 9:1 ; hexane:EtOAc to obtain cis-3-chloro-[4-(benzoxazol-2-yl)benzyloxy]-cyclopentane which is used directly in the next step.

**Step C:**     Trans-3-azido-[4-(benzoxazol-2-yl)benzyloxy]cyclopentane

5

A mixture of cis-3-chloro-[4-(benzoxazol-2-yl)benzyloxy]cyclopentane (100 mg; 0.33 ml), sodium azide (44 mg; 0.67 mmol) in 5 ml dry DMF is heated at 80°C for 24 hrs. This mixture is then concentrated, poured into water, extracted with 2 x 100 ml ether, dried (Na<sub>2</sub>SO<sub>4</sub>), concentrated in vacuo to obtain trans-3-azido-[4-(benzoxazol-2-yl)benzyloxy]cyclopentane and used directly in the next step.

10

**Step D:**     Trans-3-amino-[4-(benzoxazol-2-yl)benzyloxy]cyclopentane hydrochloride

15

A mixture of trans-3-azido-[4-(benzoxazol-2-yl)benzyloxy]cyclopentane (110 mg) and 10% Pd/C (20 mg) in 30 ml of dry ethanol is submitted to Paar hydrogenation for 2 hrs. at 40 p.s.i. and room temperature. This mixture is then filtered through celite and washed with 1 x 100 ml dry ethanol and concentrated to obtain a white solid which is dissolved in ether and precipitated as the hydrochloride salt with ethanolic HCl, filtered, washed with cold ether, dried in vacuo to obtain trans-3-amino-[4-(benzoxazol-2-yl)benzyloxy]cyclopentane hydrochloride (m.p. 235°C).

20

25

### Example 6

1-amino-[4-(benzoxazol-2-yl)benzyloxymethyl]cyclopentane hydrochloride

30

**Step A:**     1-hydroxymethyl-1-triphenylmethylaminocyclopentane

A mixture of 1-amino-1-cyclopentanemethanol (3.00 g; 26.05 mmol) and triphenylmethylchloride (7.26 g; 26.05 mmol) in 70 ml CH<sub>2</sub>Cl<sub>2</sub> is heated overnight. The reaction mixture is cooled and flash chromatographed twice using 5:1 ; hexane:EtOAc to obtain 1-hydroxymethyl-1-triphenylmethylamino-cyclopentane as a thick yellow oil which is used directly in the next step.

35



**Step B:** 1-triphenylmethylamino-(4-{benzoxazol-2-yl}benzyloxymethyl)-cyclopentane

5 To a mixture of 1-hydroxymethyl-1-triphenylmethylaminocyclopentane (2.95 g; 8.25 mmol) in 100 ml of THF is added sodium hydride (60%; 0.495 g; 12.38 mmol) and allowed to stir for 1 hr. To this mixture is then added 4-(benzoxazol-2-yl)benzyl bromide (1.90 g; 6.60 mmol) and heated to reflux for 48 hrs. The reaction mixture is quenched with H<sub>2</sub>O, concentrated in vacuo and flash  
10 chromatographed using 7:1 ; hexane:EtOAc to obtain 1-triphenylmethylamino-(4-[benzoxazol-2-yl]benzyloxymethyl)cyclopentane which is used directly in the next step.

**Step C:** 1-amino-[4-(benzoxazol-2-yl)benzyloxymethyl]cyclopentane  
15 hydrochloride

To a solution of 1-triphenylmethylamino-(4-[benzoxazol-2-yl]benzyloxymethyl)cyclopentane (1.24 g; 2.20 mmol) of 1-triphenylmethylamino-(4-[benzoxazol-2-yl]benzyloxymethyl)cyclopentane in ethanol/CH<sub>2</sub>Cl<sub>2</sub> is added ethanolic  
20 HCl until pH $\approx$ 2. This is then stirred for 72 hrs., concentrated to about one-half the volume and the white precipitate which forms is filtered to obtain 1-amino-[4-(benzoxazol-2-yl)benzyloxymethyl]cyclopentane hydrochloride (m.p. >250°C).

25 **Example 7**

2-(2-[4-(benzoxazol-2-yl)phenyl] trans-ethenyl) trans-cyclopropylformic acid

2-(2-[4-(benzoxazol-2-yl)phenyl] cis-ethenyl) trans-cyclopropylformic acid

30

**Step A:** 4-(benzoxazol-2-yl)benzyltriphenylphosphonium bromide

A mixture of 4-(benzoxazol-2-yl)benzyl bromide (0.6 g; 2.08 mmol) and triphenylphosphine (0.55 g; 2.08 mmol) in 40 ml benzene is refluxed for 48 hrs.  
35 The reaction mixture is cooled, the white solid which precipitates is filtered off, dried under vacuum to get 4-(benzoxazol-2-yl)benzyltriphenylphosphonium bromide and used directly in the next step.

**Step B:** ethyl-2-(2-[4-(benzoxazol-2-yl)phenyl] trans-ethenyl) trans-cyclopropylformate

5 ethyl-2-(2-[4-(benzoxazol-2-yl)phenyl] cis-ethenyl) trans-cyclopropylformate

To a mixture of 4-(benzoxazol-2-yl)benzyltriphenylphosphonium bromide (0.5 g; 0.908 mmol) in 15 ml DMF is added sodium hydride (60%; 0.073 g; 1.82 mmol). To this, after about 30 min., is added ethyl 2-formyl-1-cyclopropane-carboxylate (0.132 ml; 1 mmol) dropwise. The color changes from orange to yellow in about 5 min. This is stirred overnight, quenched with H<sub>2</sub>O, concentrated in vacuo and flash chromatographed using hexane:ethylacetate ; 9:1 which separates the cis (clear oil) and trans (white solid) isomers of ethyl 2-(2-[4-(benzoxazol-2-yl)phenyl]-trans-ethenyl) trans-cyclopropylformate which are confirmed by NMR before using in the next step.

**Step C:** 2-(2-[4-(benzoxazol-2-yl)phenyl] cis-ethenyl) trans-cyclopropylformic acid

20

To a mixture of ethyl 2-(2-[4-(benzoxazol-2-yl)phenyl] trans-ethenyl) trans-cyclopropylformate (0.115 g; 0.345 mmol) in 50 ml of ethanol:water ; 4:1 is added lithium hydroxide monohydrate (0.017 g; 0.414 mmol) and heated at 40°C for 7 hrs. To this is added another 1/2 equivalent of lithium hydroxide monohydrate and heating continued at 40°C overnight. This reaction mixture is then cooled, concentrated in vacuo, dissolved in dil. NaOH, extracted with ethylacetate and acidified with aqueous HCl to form a white solid. The latter is filtered and dried overnight to obtain 2-(2-[4-(benzoxazol-2-yl)phenyl] cis-ethenyl) trans-cyclopropyl-formic acid (m.p. 171-172°C).

30

**Step D:** 2-(2-[4-(benzoxazol-2-yl)phenyl] trans-ethenyl) trans-cyclopropylformic acid

When the procedure of Step C is followed using ethyl-2-(2-[4-(benzoxazol-2-yl)phenyl] cis-ethenyl) trans-cyclopropylformate the product obtained is 2-(2-[4-(benzoxazol-2-yl)phenyl] trans-ethenyl) trans-cyclopropylformic acid (m.p. 199-200°C).

35

**Example 8**5      **Trans-4-(4-[benzoxazol-yl]benzyloxy)cyclohexylamine****Step A:**      **Trans-4-triphenylmethylaminocyclohexanol**

10      A mixture of trans-4-aminocyclohexanol hydrochloride (2.00 g; 13.19 mmol), triphenylmethyl chloride (4.41 g; 15.83 mmol) and triethylamine (6.44 ml; 46.17 mmol) in 40 ml CH<sub>2</sub>Cl<sub>2</sub> is refluxed overnight. The solids are filtered off and the residue concentrated in vacuo. The resultant product is purified with CH<sub>2</sub>Cl<sub>2</sub> ⇒ CH<sub>2</sub>Cl<sub>2</sub> : ethylacetate (5:1) to obtain trans-4-triphenylmethylaminocyclo-hexanol which is used directly in the next step.

15

**Step B:**      **Trans-4-triphenylmethylamino[4-(benzoxazol-2-yl)benzyloxy]-cyclohexane**

20      To a mixture of trans-4-triphenylmethylaminocyclohexanol (3.18 g; 8.90 mmol) is 100 ml THF is added sodium hydride (60%; 0.320 g; 13.35 mmol) and this is followed by the addition of 4-(benzoxazol-2-yl)benzyl bromide after about 1 hr. The mixture is heated at reflux for 1 week, cooled, concentrated in vacuo and dissolved in about 40 ml CH<sub>2</sub>Cl<sub>2</sub> ; hexane:ethylacetate (9:1). The solid material is filtered off and the residue flash chromatographed using  
25      hexane:ethylacetate ; 7:1 to obtain trans-4-triphenylmethylamino-[4-(benzoxazol-2-yl)benzyloxy]cyclohexane as a white solid which is used directly in the next step.

30      **Step C:**      **Trans-4-[4-(benzoxazol-2-yl)benzyloxy]cyclohexanol amine hydrochloride**

To a mixture of trans-4-triphenylmethylamino-[4-(benzoxazol-2-yl)benzyloxy]-cyclohexane (0.316 g; 0.578 mmol) in 75 ml ethanol is added ethanolic HCl to pH≅2. This is stirred overnight, cooled, concentrated in vacuo  
35      and flash chromatographed using CH<sub>2</sub>Cl<sub>2</sub>:EtOH ; 3:1 to obtain trans-4-[4-(benzoxazol-2-yl)benzyloxy]cyclohexanol amine hydrochloride which is confirmed by NMR. (m.p. >250°C).

40      **Example 9**

Trans-2-(4-[benzoxazol-2-yl]benzyloxy)cycloheptylamine hydrochloride

**Step A:** Cycloheptene oxide

5

To a cooled mixture of m-chloroperbenzoic acid (9.87 g; 57.19 mmol) in 125 ml CH<sub>2</sub>Cl<sub>2</sub> is slowly added cycloheptene (5.00 g; 51.99 mmol). The reaction mixture is gradually allowed to warm to room temperature overnight. The mixture is filtered, wash with concentrated NaHCO<sub>3</sub> and concentrated in vacuo to obtain cycloheptene oxide as a pale yellow oil which is used directly in the next step.

**Step B:** Trans-2-aminocycloheptanol

15

In a sealed tube, cycloheptene oxide (5.44 g; 48.50 mmol) and ammonium hydroxide solution (10 ml) in 4 ml methanol is heated to 80°C for 72 hrs. After removal of the heat, the mixture is concentrated in vacuo to obtain trans-2-aminocycloheptanol which is used directly in the next step.

20

**Step C:** Trans-2-triphenylmethylaminocycloheptanol

A mixture of trans-2-aminocycloheptanol (0.97 g; 7.74 mmol), triphenylmethyl chloride (2.16 g; 7.74 mmol) and triethylamine (2.70 ml; 19.35 mmol) are combined in about 100 ml CH<sub>2</sub>Cl<sub>2</sub> and heated to reflux overnight. After removing the heat, the reaction mixture is concentrated in vacuo and flash chromatographed using hexane:ethylacetate; 9:1 to obtain trans-2-triphenylmethylaminocycloheptanol which is used directly in the next step.

30

**Step D:** Trans-2-triphenylmethylamino-[4-(benzoxazol-2-yl)benzyloxy]cycloheptanol

To a mixture of trans-2-triphenylmethylaminocycloheptanol (0.71 g; 1.91 mmol) in 30 ml THF and 5 ml DMF is added sodium hydride (60%; 0.115g; 2.87 mmol). After 45 min. is added 4-(benzoxazol-2-yl)benzyl bromide and potassium iodide (pinch). The reaction mixture is stirred overnight, quenched with H<sub>2</sub>O, concentrated in vacuo and flash chromatographed using exane:ethylacetate; 9:1 to obtain trans-2-triphenylmethylamino-[4-(benzoxazol-

2-yl)benzyloxy]cycloheptanol as a white solid which is used directly in the next step.

5      **Step E:**      Trans-2-(4-[benzoxazol-2-yl]benzyloxy)cycloheptylamine hydrochloride

10      To a mixture of trans-2-triphenylmethylamino-[4-(benzoxazol-2-yl)benzyloxy]cycloheptanol (0.454 g; 0.784 mmol) in 50 ml ether is added ethanolic HCl. This is stirred overnight, the white precipitate is filtered off. The latter indicates the presence of some starting material by NMR and is redissolved in EtOH/CH<sub>2</sub>Cl<sub>2</sub> and acidified to pH $\approx$ 2. This is again stirred overnight, concentrated in vacuo and flash chromatographed using CH<sub>2</sub>Cl<sub>2</sub>:ethanol; 5:1 to obtain trans-2-(4-[benzoxazol-2-yl]benzyloxy)cycloheptylamine hydrochloride (m.p. 249-250°C dec).

15

**Example 10**

20      Trans-2-aminomethyl-[4-(benzoxazol-2-yl)benzyloxymethyl]cyclopropane hydrochloride

**Step A:**      Ethyl 2-triphenylmethylaminomethylcyclopropane-1-carboxylate

25      To a mixture of ethyl 2-formylcyclopropane-1-carboxylate (1.00 g; 7.03 mmol) and magnesium sulfate (0.85g; 7.03 mmol) in 50 ml methanol is added triethylamine (1.82 g; 7.03 mmol). To this is then added sodium cyanoborohydride (0.88 g; 14.06 mmol) and stirred for 48 hrs. To this is then further added 0.20 g of ethyl 2-formylcyclopropane-1-carboxylate and stirring continued for another 72 hrs. The reaction mixture is filtered, concentrated in vacuo and flash chromatographed using hexane:ethylacetate ; 9:1 to obtain ethyl-2-triphenylmethylaminomethyl-cyclopropane-1-carboxylate as a colorless oil which is used directly in the next step.

30

**Step B:**      2-triphenylmethylaminomethyl-1-hydroxymethylcyclopropane

35

To a mixture of lithium aluminum hydride (0.077 g; 2.04 mmol) in 10 ml THF is added ethyl 2-triphenylmethylaminomethylcyclopropane-1-carboxylate (0.714 g; 1.85 mmol) in 20 ml THF and stirred overnight. The reaction mixture is quenched with water, filtered, concentrated in vacuo and flash chromatographed

using hexane:ethylacetate ; 3:1 to obtain 2-triphenylmethylaminomethyl-1-hydroxy-methylcyclopropane as a clear oil which is used directly in the next step.

5 **Step C:** Trans-2-triphenylmethylaminomethyl-[4-(benzoxazol-2-yl)benzyloxymethyl]cyclopropane

To a mixture of 2-triphenylmethylaminomethyl-1-hydroxymethyl-cyclopropane (0.338 g; 0.984 mmol) in 30 ml THF and 5 ml DMF is added  
10 sodium hydride (60%; 0.059 g; 1.48 mmol) followed by 4-(benzoxazol-2-yl)benzyl bromide (0.255 g; 0.886 mmol) and a pinch of potassium iodide. This is stirred overnight, quenched with water, concentrated in vacuo and flash chromatographed using hexane:ethyl-acetate ; 7:1 to obtain trans-2-  
15 triphenylmethylaminomethyl-[4-(benzoxazol-2-yl)-benzyloxymethyl]cyclopropane as a pale yellow oil which is used directly in the next step.

**Step D:** Trans-2-aminomethyl-[4-(benzoxazol-2-yl)benzyloxymethyl]cyclopropane hydrochloride

20 To a mixture of trans-2-triphenylmethylaminomethyl-[4-(benzoxazol-2-yl)benzyloxymethyl]cyclopropane (0.252 g; 0.458 mmol) in 40 ml EtOH:CH<sub>2</sub>Cl<sub>2</sub> ; 10:1 is added ethanolic HCl until pH=2. This is stirred overnight, further ethanolic HCl is added and stirring continued another 24 hrs. The reaction mixture is then heated to reflux for 5 hrs., concentrated in vacuo and flash  
25 chromatographed using CH<sub>2</sub>Cl<sub>2</sub>: EtOH ; 3:1 to obtain trans-2-aminomethyl-[4-(benzoxazol-2-yl)benzyloxy-methyl]cyclopropane hydrochloride as a white solid (m.p. >250°C).

30 **Example 11**

Trans-2-[4-(benzoxazol-2-yl)benzyloxy]cyclopentylamine acetate

**Step A:** Trans-2-aminocyclopentanol

35

A mixture of cyclopentene oxide (5.00 g ; 59.44 mmol) ammonium hydroxide solution (10 ml) and 4 ml methanol are placed in a sealed tube and heated at 80°C for 72 hrs. The reaction mixture is then concentrated in vacuo and flash chromatographed using CH<sub>2</sub>Cl<sub>2</sub>:EtOH ; 1:1 to obtain trans-2-  
40 aminocyclopentanol which is used directly in the next step.

**Step B:**     Trans-N-(2-hydroxy)cyclopropylphthalimide

5         A mixture of trans-2-aminocyclopentanol (6.00 g; 59.32 mmol), phthalic anhydride (10.54 g; 71.18 mmol) and triethylamine (28.94 ml; 207.62 mmol) in 80 ml CHCl<sub>3</sub> are heated at reflux for 72 hrs. The reaction mixture is then concentrated in vacuo and flash chromatographed using hexane:ethylacetate ; 1:1 to obtain trans-N-(2-hydroxy)cyclopropylphthalimide which is used directly in the next step.

10

**Step C:**     Trans-2-(N-phthalimido)-1-[4-(benzoxazol-2-yl)benzyloxy]cyclopentane

15         To a mixture of sodium hydride (60%; 0.322 g; 8.06 mmol) in 15 ml THF at 0°C is added trans-N-(2-hydroxy)cyclopropylphthalimide (1.434 g; 6.20 mmol) in 10 ml THF. After about 1 hr., 4-(benzoxazol-2-yl)benzyl bromide (1.965 g; 6.82 mmol) in 5 ml DMF is added. The reaction mixture is stirred overnight, quenched with water, concentrated in vacuo and flash chromatographed using hexane:ethyl-acetate ; 3:1 to obtain trans-2-(N-phthalimido)-1-[4-(benzoxazol-2-yl)benzyl-oxy]cyclopentane as a white solid which is used directly in the next step.

20

**Step D:**     Trans-2-[4-(benzoxazol-2-yl)benzyloxy]cyclopentylamine acetate

25         To a mixture of trans-2-(N-phthalimido)-1-[4-(benzoxazol-2-yl)benzyloxy]cyclopentane (0.400 g; 0.912 mmol) in 1.4 ml water and 8 ml 2-propanol is added sodium borohydride (0.173 g; 4.56 mmol) and stirred overnight, concentrated in vacuo and flash chromatographed using CH<sub>2</sub>Cl<sub>2</sub>:EtOH; 20:1 ⇒ CH<sub>2</sub>Cl<sub>2</sub>:EtOH; 15:1 to obtain an off-white solid. This material is added to 8 ml of 2-propanol and 1.4 ml water to which is added 1 ml acetic acid and heated to 80°C for 2 hrs. Upon concentration in vacuo and flash chromatography using CH<sub>2</sub>Cl<sub>2</sub>:EtOH ; 5:1 trans-2-[4-(benzoxazol-2-yl)benzyloxy]cyclopentylamine acetate is obtained (m.p. >200°C).

30

35

**Example 12**

2 $\beta$ -hydroxy-3 $\beta$ [(4-benzoxazol-2-yl)benzyloxy]cyclohexyl-1 $\alpha$ -amine

**Step A:**     Syn-2,3-epoxycyclohexanol

5           To 2-cyclohexan-1-ol (5.0 g, 51 mmol) in 100 mL dry dichloromethane at 0°C under argon is added in portions m-chloroperbenzoic acid (13.2 g, 61 mmol) over five minutes. The mixture is permitted to warm slowly to room temperature and is stirred overnight. Residual solid is filtered off and the supernatant washed with 2 x 100 mL portions of saturated sodium bicarbonate solution. The  
10 organics are dried (sodium sulfate) and concentrated. Flash chromatography (gradient elution ; dichloromethane to 10% methanol : dichloromethane) affords syn-2,3-epoxycyclohexanol which is further purified by distillation [Tetrahedron Lett. 4895 (1985)] and used directly in the next step.

15   **Step B:**     Syn-2-(4'-benzoxazol-2"-ylbenzyloxy)cyclohexane epoxide

          To potassium hydride (2.1 g, 18.4 mmol, 35%) in 50 mL dry THF at 0°C is added syn-2-epoxycyclohexan-1-ol (1.91 g, 16.7 mmol). The mixture is stirred at 0°C for two hours at which point 4-benzoxazolebenzylbromide (4.81 g,  
20 16.7 mmol) is added in one portion. The mixture is permitted to warm to room temperature and is stirred for one hour. The reaction is quenched by the addition of water and concentrated. Flash chromatography (gradient elution 6:1  $\Rightarrow$  3:1 ; hexane:ethyl acetate) affords syn-2-(4'-benzoxazol-2"-ylbenzyloxy)-cyclohexane epoxide which is used directly in the next step.

25

**Step C:**     2 $\beta$ -hydroxy-3 $\beta$ [(4-benzoxazol-2-yl)benzyloxy]cyclohexyl-1 $\alpha$ -amine

          2-(4'-benzoxazol-2"-ylbenzyloxy)cyclohexane epoxide (4.22 g) is combined in a sealed tube with concentrated ammonium hydroxide solution (5 mL) and methanol (20 mL) and heated at 80°C for 12 hours. The mixture is  
30 concentrated and subjected to flash chromatography (gradient elution methanol  $\Rightarrow$  5% triethylamine : methanol) to afford a white solid which is recrystallized from ethanol to obtain 2 $\beta$ -hydroxy-3 $\beta$ [(4-benzoxazol-2-yl)benzyloxy]cyclohexyl-1 $\alpha$ -amine. (m.p.149°C).

35

**Example 13**



When the above procedures are followed but the appropriate starting materials are used, then the desired compound may be obtained. A representative list of compounds which may be prepared include the following:

- 5  
 6- $\alpha$ -hydroxy-2 $\alpha$ -[(4-benzoxazol-2-yl)benzyloxy]-cyclohexyl-1 $\beta$ -amine  
 hydrochloride (m. p. 242-248°C)  
trans-2-(4-[2-benzoxazolyl]-benzyloxy)cyclobutylamine hydrochloride  
 (m. p. 151-155°C)  
 10 trans-2-(4-[4-chlorobenzoyl]-benzyloxy)cyclohexylamine hydrochloride  
 (Exact mass: (M+H)<sup>+</sup>:calc.=344.1417 ; found=344.1400)  
cis-2-(4-[2-benzoxazolyl]-benzyloxy)cyclobutylamine hydrochloride  
 (m. p. 250-251°C)  
trans-2-(4-[2-benzoxazolyl]-benzyloxy)-1,2,3,4-tetrahydre-1-naphthylamine  
 15 hydrochloride (m. p. >250°C)  
trans-2-(4-[2-benzoxazolyl]-benzyloxymethyl)cyclopropyl-methylamine  
 hydrochloride (m. p. >250°C)  
 1 $\beta$ ,2 $\beta$ -dihydroxy-1 $\alpha$ -[4-(benzoxazol-2-yl)phenyl]  
 cyclohexyl-3 $\alpha$ -amine (m.p. 183-185°C)  
 20 2 $\beta$ -hydroxy-1 $\beta$ -[4-(benzoxazol-2-yl)benzyloxy]cyclohexyl-4 $\alpha$ -amine  
 4 $\beta$ -hydroxy-1 $\beta$ -[4-(benzoxazol-2-yl)benzyloxy]cyclohexyl-3 $\alpha$ -amine  
 2 $\beta$ -hydroxy-1 $\beta$ -[4-(benzoxazol-2-yl)benzyloxy]cyclopentyl-3 $\alpha$ -amine  
 2 $\beta$ -hydroxy-1 $\beta$ -[4-(benzoxazol-2-yl)benzyloxy]cycloheptyl-3 $\alpha$ -amine  
 2 $\alpha$ -hydroxy-1 $\beta$ -[4-(benzoxazol-2-yl)benzyloxy]cyclopentyl-3 $\beta$ -amine  
 25

#### Example 14

When the above procedures are followed but the appropriate starting materials are used, then the desired compound may be obtained. A representative list of compounds which may be prepared include the following:

- 2-[4-(benzoxazol-2-yl)benzyloxy]cyclohexylamine  
 3-[4-(benzoxazol-2-yl)benzyloxy]cyclohexylamine  
 2-[4-(benzthiazol-2-yl)benzyloxy]cyclohexylamine  
 2-[3-(benzthiazol-2-yl)benzyloxy]cyclohexylamine  
 2-[4-(benzthiazol-2-yl)benzyloxy]cyclopentylamine  
 2-[4-benzoylbenzoxy]cyclohexylamine  
 35

- 2-[4-(3,4-methylenedioxy)benzyloxy]cyclohexylamine  
 2-[4-(2H-3,4-dihydro-1-benzopyran-6-yl)benzyloxy]cyclohexylamine  
 2-[4-(2,3-dihydrobenzofuran-5-yl)benzyloxy]cyclohexylamine  
 2-[4-(2,3-dihydro-1,4-benzodioxin-6-yl)benzyloxy]cyclohexylamine  
 5 2-[4-benzo[b]thien-2-yl)benzyloxy]cyclohexylamine  
 4-[4-(quinolin-2-yl)benzyloxy]cyclohexylamine  
 2-[4-(quinolin-2-yl)benzyloxy]cyclohexylamine  
 2-[4-(quinolin-2-yl)benzyloxy]cyclopentylamine  
 4-[4-naph-2-yl)benzyloxy]cyclohexylamine  
 10 4-[4-naph-1-yl)benzyloxy]cyclohexylamine  
 2-[4-naph-1-yl)benzyloxy]cyclohexylamine  
 4-[4-(1-methylindol-2-yl)benzyloxy]cyclohexylamine  
 2-[4-benzofuran-2-yl)benzyloxy]cyclohexylamine  
 4-[4-(3-methoxyphenyl)benzyloxy]cyclohexylamine  
 15 4-[4-(3-methoxyphenyl)benzylthio]cyclohexylamine  
 2-[4-(2-(E)phenylethenyl)benzyloxy]cyclohexylamine  
 2-[4-(benzoxazol-2-yl)naph-1-yl]methoxycyclohexylamine  
 3-[5-(benzoxazol-2-yl)thien-2-yl]methoxycyclohexylamine  
 2-[5-(benzoxazol-2-yl)thien-2-yl]methoxycyclohexylamine  
 20 2-[4-(benzoxazol-2-yl)phenoxy]cyclohexylamine  
 2-[4-(benzoxazol-2-yl)phenethoxy]cyclohexylamine  
 4-[4-(benzthiazol-2-yl)benzylthio]cyclohexylamine  
 4-[4-(benzoylphenyl)phenethyl]cyclohexylamine  
 4-[4-(benzoylphenyl)benzyl]cyclohexylamine  
 25

### Example 15

When the above procedures are followed but the appropriate starting  
 30 materials are used, then the desired compound may be obtained. A  
 representative list of compounds which may be prepared include the following:

- 3- $\beta$ -(4-[benzofuran-2-yl]-benzyloxy)-2- $\beta$ -hydroxy-cyclohexyl- $\alpha$ -amine  
 hydrochloride [m. p. 208-212°C (dec)]  
 35  
 2- $\alpha$ -hydroxy-2 $\beta$ -methyl-3 $\beta$ -[(4'-benzoxazol-2''-yl)benzyloxy]cyclohexyl-1 $\beta$ -amine  
 (m. p. 118-119°C)

- 3- $\beta$ -(4-[2-benzothiazolyl]benzyloxy)-2- $\beta$ -hydroxy-cyclopentyl- $\alpha$ -amine  
[m. p. 192-195°C (dec.)]
- 5      2- $\beta$ -[4-(benzoxazol-2'-yl)phenyl]-6- $\beta$ -hydroxy-6- $\alpha$ -aminehydrochloride  
(214-216°C)
- (1 $\alpha$ ,2 $\alpha$ )-(+-)-2-[4-(benzoxazol-2-yl)phenyl]2-hydroxy-1-cyclohexylamine  
hydrochloride [m. p. 170°C (dec)]
- 10      2- $\beta$ -[(4'- $\beta$ ' - naphthyl)benzyloxy]-6- $\beta$ -hydroxy-6- $\alpha$ -methylcyclo-hexyl-1- $\alpha$ -amine  
hydrochloride (m. p. 136-138°C)
- 15      3- $\beta$ -(4-[2-naphthyl]benzyloxy)-2- $\beta$ -hydroxycyclohexyl- $\alpha$ -amine hydrochloride  
(m. p. 207-209°C)
- 3- $\beta$ -(4-[2-benzothiazolyl]-benzyloxy)-2- $\beta$ -hydroxycyclohexyl- $\alpha$ -amine  
dihydrochloride (m. p. 135-139°C)
- 20      3- $\alpha$ -(4-[2-benzoxazolyl]-phenyl)-2- $\beta$ -hydroxycyclo-hexyl- $\alpha$ -amine hydrochloride  
(m. p. >250°C)
- 2- $\alpha$ -(4-[benzoxazol-2-yl]-phenyl)-2- $\beta$ -hydroxycyclohexyl- $\alpha$ -amine hydrochloride  
(m. p. >250°C)
- 25      2- $\alpha$ -hydroxy-3 $\beta$ -[(4'-benzoxazol-2''-yl)naphthylene-1-methyloxy]cyclohexyl-  
1 $\beta$ -amine dihydrochloride (m. p. 142-145°C)
- (-)-2 $\alpha$ -hydroxy-3 $\beta$ -[(4'-benzoxazol-2''-yl)benzyloxy]-cyclohexyl-1 $\beta$ -amine
- 30      (+)-2 $\alpha$ -hydroxy-3 $\beta$ -[(4'-benzoxazol-2''-yl)benzyloxy]-cyclohexyl-1 $\beta$ -amine
- N,N-dimethyl-2- $\alpha$ -hydroxy-2 $\beta$ -methyl-3 $\beta$ -[(4'-benzoxazol-2''-yl)benzyloxy]-  
cyclohexyl-1 $\beta$ -amine hydrochloride (m. p. 241-242°C)
- 35      N,N-dimethyl-3- $\beta$ -(4-[2-benzoxazolyl]benzyloxy)-2- $\beta$ -hydroxycyclohexyl- $\alpha$ -amine  
hydrochloride (m. p. 76-79°C)

- 2- $\alpha$ -methoxy-3 $\beta$ -[4'-benzoxazol-2''-yl]benzyloxy]cyclo-hexyl-1  $\beta$ -amine  
hydrochloride (m. p. 270°C)
- 5 [1 $\alpha$ ,2 $\alpha$ ,3 $\beta$ (+ -)]3-[4-(2-benzthiazolyl)phenoxy]-2-hydroxy-1-cyclohexylamine  
(m.p. 154-158°C)
- [1 $\alpha$ ,2 $\alpha$ ,3 $\beta$ (+ -)]3-[4-(2-benzothiazolyl)phenoxy]-2-hydroxy-1-cyclohexylamine  
(m.p. 170-172°C)
- 10 [1 $\alpha$ ,2 $\beta$ ,3 $\beta$ (+ -)]2-hydroxy-3[4-(2-quinoliny)]phenylmethoxy]-1-cyclohexylamine  
dioxalic acid (m.p. 199-201°C)
- [1 $\alpha$ ,2 $\beta$ ,3 $\beta$ (+ -)]2-hydroxy-3[4-(3-methoxyphenyl)phenyl-methoxy]-1-  
cyclohexylamine hemioxalic acid [m.p. 233°C (dec)]
- 15 (1  $\alpha$ ,2 $\beta$ ,3 $\alpha$ )-( + -)-3-[4-(benzoxazol-2-yl)]phenylmethyl]-2-hydroxy-1-  
cyclohexylamine (m.p. 164-166°C)
- (1  $\alpha$ ,2 $\alpha$ ,3 $\alpha$ )-( + -)-3-[4-(benzoxazol-2-yl)]phenylmethyl]-2-hydroxy-1-  
cyclohexylamine hydrochloride [m.p. 285°C (dec)]
- 20 (1  $\alpha$ ,2 $\alpha$ ,3 $\beta$ )-( + -)-2-[4-(benzoxazol-2-yl)]phenyl]-2-hydroxy-3-methyl-1-  
cyclohexylamine acetate (m.p. 187-195°C)
- 25 (1  $\alpha$ ,2 $\alpha$ ,3 $\beta$ )-( + -)-2-[4-(benzoxazol-2-yl)]phenyl]-3-chloro-2-hydroxy-1-  
cyclohexylamine hydrochloride [m.p. 275°C (dec)]
- (1  $\alpha$ ,2 $\beta$ ,3 $\beta$ )-( + -)-2-[4-(benzoxazol-2-yl)]phenyl]-2,3-dihydroxy-1-cyclohexylamine  
hydrochloride [m.p. 285°C (dec)]
- 30 (1  $\alpha$ ,2 $\alpha$ ,3 $\alpha$ )-( + -)-2-[4-(benzoxazol-2-yl)]phenyl]-2,3-dihydroxy-1-cyclohexylamine  
hydrochloride [m.p. 275°C (dec)]
- (1  $\alpha$ ,2 $\alpha$ )-( + -)-2-[4-(benzoxazol-2-yl)]phenyl]-2-methoxy-1-cyclohexylamine  
hydrochloride (m.p. 190-197°C)
- 35 2-[4-(benzoxazol-2-yl)]phenyl]-cyclohex-2-enaminehydrochloride

(m.p. >310°C)

5

(1R,2R,3R)-2-[4-(benzoxazol-2-yl)phenylmethoxy]-3-(2-methoxyethyl)-1-cyclopentylamine acetate (m.p. 133-136°C)

(1R,2R,3R)-2-[4-(benzoxazol-2-yl)phenylmethoxy]-3-(2-hydroxyethyl)-1-cyclopentylamine acetate (m.p. 145-147°C)

10

(1R,2R,3S)-2-[4-(benzoxazol-2-yl)phenyl]methoxy-3-(2-methoxyethyl)-1-cyclopentylamine hydrochloride (m.p. 84-95°C)

(1R,2R,3R)-1-[4-(benzoxazol-2-yl)phenyl]methoxy-3-(2-hydroxyethyl)-2-cyclopentylamine hydrochloride (m.p. 210-213°C)

15

(1R,2R,3S)-2-[4-(benzoxazol-2-yl)phenyl]methoxy-3-(2-hydroxyethyl)-1-cyclopentylamine acetate (115-125°C)

20

trans-2-(p-(2-benzoxazolyl)phenyl)cyclohexylamine acetate  
(m.p. 162-164°C)

trans-6-(p-(2-benzoxazolyl)phenyl)cyclohex-3-enylamineacetate  
(m.p. 157-160°C)

25

(1 $\alpha$ ,2 $\beta$ ,3 $\beta$ )-(+-)-2-hydroxy-3-[4-(p-chlorobenzoyl)-phenyl-methoxy]-1-cyclohexylamine hemioxalate [m.p. 218°C (dec)]

(1R,2R,3R)-2-[4-(benzoxazol-2-yl)phenylmethoxy]-3-(2-hydroxyethyl)-1-cyclopentylamine acetate (m.p. 144-146°C)

30

(1 $\alpha$ ,2 $\alpha$ ,3 $\alpha$ )-(+-)-3-[4-(benzoxazol-2-yl)phenylmethyl]-2-hydroxy-1-cyclohexylamine hydrochloride [m.p. 285°C (dec)]

(1 $\alpha$ ,2 $\alpha$ ,3 $\alpha$ )-(+-)-2-[4-(benzoxazol-2-yl)phenyl]-2,3-dihydroxy-1-cyclohexylamine hydrochloride[m.p. 275°C (dec)]

35

(1 $\alpha$ ,2 $\beta$ ,3 $\beta$ )-(+-)-2-[4-(benzoxazol-2-yl)phenyl]-2-chloro-3-hydroxy-1-cyclohexylamine hydrochloride [m.p. 275°C (dec)]

(1 $\alpha$ ,2 $\alpha$ ,3 $\beta$ )-(+-)-2-[4-(benzoxazol-2-yl)phenyl]-2-hydroxy-3-methyl-1-cyclohexylamine acetate (m.p. 187-195°C)

5 2- $\beta$ -[4-(benzoxazol-2'-yl)benzy]-6- $\beta$ -chloro-cyclohexyl-1- $\alpha$ -amine hydrochloride  
[m. p. 236-240°C (dec)]

cis-3-(4-[2-benzoxazolyl]-benzyloxy)cyclohexylaminehydrochloride  
(m. p. 258-261°C)

10

trans-3-(4-[2-benzoxazolyl]-benzyloxy)cyclohexylaminehydrochloride  
[m. p. 240°C (dec)]

3-(4-[2-benzoxazolyl]phenyl)-cyclohexylamine hydrochloride(m. p. > 250°C)

15

Various tests have been carried out to show the ability of the compounds of the present invention to exhibit pharmacological responses that can be correlated with activity in humans. These tests involve such factors as the effect of the compounds of Formula I to inhibit squalene synthesis. It has been found that compounds within the scope of this invention when tested using the following procedures show a marked activity for the inhibition of squalene synthase and hence are believed to be useful in the treatment of cholesterol-related disorders.

25

#### Squalene Synthase Inhibition Assay

The squalene synthase assay used is a modification of the procedures described by Popjak (1969) and Poulter et al. (1989):

30

Popjak, G. Enzymes of sterol biosynthesis in liver and intermediates of sterol biosynthesis. Meth. Enzymol. 15: 393-454, 1969.

Poulter, C.D., Capson, T.L., Thompson, M.D. and Bard R.S. Squalene synthase. Inhibition by ammonium analogues of carbocationic intermediates in the conversion of presqualene diphosphate to squalene. J. Am. Chem. Soc. 111: 3734-3739, 1989.

35

I. Animal Source and Tissue Preparation:

5 Four male Sprague-Dawley rats weighing 100-120 gms are fed a low cholesterol rodent diet (#5012) obtained from Purina Mills, Inc. in Richmond, Indiana; and housed under reverse-light. Water is given ad lib. Rats are lightly anesthetized with ether and then decapitated. Livers are removed and enzymes are separated by the method described below.

10 II. Materials:

Chemicals:

All Chemicals are "A.C.S." in purity or better unless noted;

15 AquaSol® -2 scintillation fluid (NEF-952) (Du Pont/NEN Research Products, Boston, Mass.);

Anhydrous  $MgCl_2$  (M-8266),  $\beta$ -NADPH tetrasodium salt, reduced form (N-1630), Bovine serum albumin (A-6003), Cholesterol (C-8503);

20 Squalene (S-3626), (Sigma Chemical Co., St. Louis, Missouri);

Bio-Rad protein assay dye concentrate (Bio-Rad Laboratories, Richmond, CA);

25 Denatured ethanol, DMSO, HCl (1 N), KOH, methanol, NaOH (.1N, 1N), petroleum ether (M-280 grade), potassium phosphate dibasic, 2-propanol (Fisher Scientific, Pittsburgh, PA);

30 Zero grade nitrogen gas mixture (certified analysis) (Woodland Oxygen & Supply Co., Philadelphia, PA).

Radiochemicals:

35 [1- $^3H$ (N)]-FPP, triammonium salt (NET-1042), (Du Pont/NEN, Boston, Mass.);

[4,8,12,13,17,21- $^3H$ ]-Squalene (NET-645) (Du Pont/NEN);

Non-radiolabeled FPP is prepared in-house. The solid FPP is aliquoted and stored at  $-80^{\circ}\text{C}$ . FPP is dissolved in 70% ethanol/30% 0.25 M  $\text{NH}_4\text{HCO}_3$  at the concentration of 10 mM and the solution is aliquoted (200  $\mu\text{l}$  each) and  
5 stored at  $-80^{\circ}\text{C}$ .

### III. Preparation of Assay Substances:

#### A) Test Solutions:

10 Test solutions are prepared fresh in 100% DMSO or  $\text{dH}_2\text{O}$ . Subsequent dilutions are made in the same solvent. Compounds are tested initially at 1 or 10  $\mu\text{M}$  (final concentrations).

#### B) Assay Buffer:

15 Potassium Phosphate (50 mM, 8.71 g/l) pH 7.5 stock buffer is prepared and stored at  $4^{\circ}\text{C}$  until use. Anhydrous  $\text{MgCl}_2$  is added to the phosphate buffer on the day of assay for a final concentration of 10mM (95 mg/100 ml). The buffer is flushed with  $\text{N}_2$  before use.

#### 20 C) Substrate:

Non-radiolabeled FPP is diluted to 50  $\mu\text{M}$  (100  $\mu\text{l}$  10 mM cold FPP +19.9 ml phosphate buffer). Then, 14  $\mu\text{l}$  ( $20 \times 10^6$  dpm) of  $^3\text{H}$ -FPP (0.5 mCi/ml, .011 mg/ml) is added. 200  $\mu\text{l}$  of this mixture is added per assay tube for a final reaction concentration of 10  $\mu\text{M}$  FPP (~200,000 dpm/assay tube).  
25

#### D) $\beta$ -NADPH Solution:

37.5 mg of  $\beta$ -NADPH is added to 9 ml assay buffer for a 5 mM concentration of  $\beta$ -NADPH. The mixture is vortexed and 100  $\mu\text{l}$  of this solution is added to each tube for a final assay concentration of 0.5 mM  $\beta$ -NADPH.  
30

#### E) KOH in Ethanol:

75 gm of KOH is dissolved in 500 ml of denatured ethanol for a 15% solution and stored at  $0^{\circ}\text{C}$  until use. 1 ml of this solution is added per tube to terminate the reaction.  
35

### IV. Experimental Procedure:



#### A) Enzyme Preparation:

Immediately following decapitation, livers are removed one at a time from four rats. The livers are combined and weighed in a tared beaker. Assay buffer is added equal to three times the liver weight. The liver is first homogenized  
5 with a blender for thirty seconds, and then by a motor driven teflon pestle at a speed of 2.5. During homogenization, the liver is kept on ice. When the liver is fully homogenized, the homogenate is centrifuged at 10,000 g for 30 min at 4°C in 50 ml capacity centrifuge tubes. The mitochondrial pellet is discarded and the supernatant is filtered through a layer of gauze moistened with a little buffer.  
10 This supernatant is recentrifuged at 105,000 g for one hour at 0°C in an ultracentrifuge in 25 ml capacity ultracentrifuge tubes.

Following centrifugation, the supernatant is removed and discarded. The sediment pellet consists of 2 layers: a transparent inner layer of glycogen,  
15 surrounded by an opaque brown layer of microsomes. The brown outer microsomal layer is carefully removed with a spatula and placed in a beaker on ice. Assay buffer is added in an amount equal to one half the original homogenate volume, and this mixture is poured into ultracentrifuge tubes. These tubes are recentrifuged at 105,000 g for 1 hour at 4°C.

20 After this centrifugation is complete, the supernatant is again removed and discarded. Fresh assay buffer is added to the combined pellets to achieve a volume equal to one tenth of the original homogenate volume. The microsomal fraction is then rehomogenized on a motor driven teflon pestle at a  
25 speed of 2.5 to partially solubilize and make a uniform suspension of the microsomes. The enzyme (~20 ml, ~40 mg protein/ml) is aliquoted (200 µl) into eppendorf plastic tubes, capped and stored at -80°C until use.

#### B) Assay Procedure

30 To begin the assay, 20 µl of the compound of this invention or vehicle solution is added to each 16 x 150 screw-cap culture tube on ice. Then 580 µl of N<sub>2</sub> flushed assay buffer is pipetted into each tube. 100 µl of cofactor is next added to each tube, followed by 100 µl of a dilution of microsomal enzyme (approximately 80 ug protein). The tubes are preincubated for 10 minutes at  
35 37°C, and 200 µl of the <sup>3</sup>H-FPP (200,000 dpm, 10 µM final conc.) is added to each tube at two second intervals. The tubes are then incubated for exactly 10 minutes, shaking at 150 oscillations per minute. After the 10 minute incubation,

the reaction is stopped by the addition of 1 ml of 15% KOH in ethanol, and the tubes are incubated for 30 minutes in a 65°C water bath for saponification of lipids and solubilization of proteins. The tubes are cooled on ice for five minutes. The samples are next extracted with 5 ml of petroleum ether by  
5 shaking for 10 minutes at low speed on a metabolic shaker. Each lower aqueous layer is frozen in a dry ice/alcohol bath (2-propanol/methanol, 1:1), and each organic layer is poured into another set of 16 x 150 screw-top culture tubes containing 2 ml of deionized water. Each ether layer is washed by vortexing each tube for 5 seconds. The aqueous layers are again frozen in the dry  
10 ice/alcohol bath, and the ether is poured into scintillation vials. 10 ml of AquaSol® is next added to each vial, and the vials are counted for 5 minutes in a scintillation counter. Percent inhibitions are calculated from the counts obtained.

15 V. Statistical Considerations

The samples are counted as dpm using a Beckman Scintillation counter (Model LS-9000). Percent inhibition is calculated using a Lotus 1-2-3 program. The IC<sub>50</sub> values are calculated using a linear regression program of Tallarida  
20 and Murray (1987). Tallarida, R.J. and Murray, R.B. Manual of pharmacologic calculations with computer programs. Springer-Verlag, 1987.

The following in vivo assays are used to determine the effectiveness of compounds of this invention to inhibit squalene synthase.

25

In Vivo Rat Method:

Forty male Sprague-Dawley rats weighing 80-90 gms are fed a low cholesterol rodent meal diet (#5012) obtained from Purina Mills, Inc. in  
30 Richmond, Indiana; and housed under reversed-light. Water is given ad lib. After a week of housing, the rats are fed cholestyramine (2% in diet) for the next 2 days. One day after the cholestyramine treatment, a compound is given three times to 4 rats (30 mg/kg, in 10 ml 0.5% methylcellulose/Kg, p.o.) at 8 a.m., 5 p.m. and 8 a.m. the following day. Four hours after the last dose, the rat is  
35 decapitated and blood and the liver are collected. Serum is separated and analyzed for cholesterol by an assay kit supplied by Sigma Chemical Co., St. Louis, Missouri. Sterols in the serum and the liver are also analyzed by HPLC

analysis. The liver cholesterol level is determined by quantitating against cholesterol as a standard by HPLC. Results are determined as the change in the liver and serum cholesterol compared to respective vehicle treated control values.

5

Mouse in vivo assay for cholesterol-lowering effect:

Balb-c mice weighing 20-25 gms are used for this study. Mice are fed a meal diet (RP#5012) containing 2% cholestyramine. Test compound is dissolved/suspended in 0.5% methyl cellulose and given to mice at 50 mg/kg, b.i.d. (n=8). Blood is collected from the tail on days 0 and 7 and from inf. vena cava on day 14. Body weights and the food consumption are monitored weekly. Serum cholesterol is determined by an enzyme assay kit (Sigma Chemical Co.). Serum and the liver are analyzed by HPLC for various sterols. Results are determined as % change from the vehicle treated controls.

HPLC Procedure

Sample preparation:

20

In 16x125 mm screw cap glass tubes, is added 0.5 ml serum or plasma or liver slices suspension (0.11 gm/ml buffer). To this is then added 1 ml of KOH/ethanol (15%). The glass tubes are incubated at 80°C for 2 hours while shaking. After saponification, 10 ml petroleum ether is added to each tube and then capped. The tubes are shaken for 30 min. then frozen at -80°C and the ether removed to 16x150 mm screw top glass tubes. Water (2 ml ) is added and again the tubes are shaken for 10 minutes. After freezing, the ether is transferred into 16x125 mm glass culture tubes. The ether is evaporatedf under N<sub>2</sub>. HPLC grade ethanol (0.5 ml) is added to the tubes, capped and shaken for 30 min to dissolve all lipids. The ethanol is filtered using a syringe filter (pore size:0.45 uM, Spartan-13, Schleicher & Schuell) and 50 ul of this solution is injected into HPLC.

25  
30

HPLC specifications:

35

Column: Partisil 10 ODS-3, C-18 reversed phase, 25 cm length at 25°C.

[Whatman Cat#4228-001]

- Pump: Waters 6000A, Isocratic system  
Solvent: Acetonitrile:water:: 95:5; 2 ml/min  
Run Time: 33 min  
50 ul injection by Autosampler WISP 710B, Waters/Millipore  
5 Wavelength: 210 nm, Applied Biosystems 757 Absorbance detector  
Integrator: Hewlett Packard 3396 Series II.

Typical Retention Times:

- 10 Squalene dioxide - 7.5 min  
Squalene oxide - 14 min  
7-dehydro Chol. - 16 min  
Cholesterol - 23 min  
Squalene - 30 min

15

- Compounds within the scope of Formula I have been tested by the foregoing assay procedures and exhibit marked squalene synthase inhibition activity and are useful as hypocholesterolemic or hypolipidemic agents by virtue of their ability to inhibit the biosynthesis of cholesterol. Having such ability, the compounds are incorporated into pharmaceutically acceptable carriers and administered to a patient in need of such cholesterol biosynthesis inhibition. These pharmaceutical formulations contain at least one compound according to this invention.

- 25 Treatment with a combination of an HMG-CoA reductase inhibitor and a squalene synthase inhibitor would have a synergistic effect on inhibiting cholesterol biosynthesis. Inhibiting the squalene synthase enzyme and the HMG-CoA reductase enzyme at the same time would most closely resemble the physiological conditions of cholesterol homeostasis. A squalene synthase inhibitor could keep cellular concentrations of farnesyl diphosphate high enough for the synthesis of the small amounts of dolichol, ubiquinone, and the farnesylated proteins required by the cell. This would maintain some feedback regulation of the HMG-CoA reductase enzyme and allow smaller amounts of the HMG-CoA reductase inhibitor to be used.

35

Other combinations with a squalene synthase inhibitor which could have a synergistic effect for controlling undesirable cholesterol levels in the body

include niacin, antihyperlipoproteinemic agents such as gemfibrozil, cholesterol absorption inhibitors, bile acid sequestrants, antioxidants and lipoxigenase inhibitors.

5           The compounds of the present invention can be administered to a mammalian host in a variety of forms adapted to the chosen route of administration, i.e., orally, or parenterally. Parenteral administration in this respect includes administration by the following routes: intravenous, intramuscular, subcutaneous, intraocular, intrasynovial, transepithelially  
10 including transdermal, ophthalmic, sublingual and buccal; topically including ophthalmic, dermal, ocular, rectal and nasal inhalation via insufflation and aerosol and rectal systemic.

          The active compound may be orally administered, for example, with an  
15 inert diluent or with an assimilable edible carrier, or it may be enclosed in hard or soft shell gelatin capsules, or it may be compressed into tablets, or it may be incorporated directly with the food of the diet. For oral therapeutic administration, the active compound may be incorporated with excipient and used in the form of ingestible tablets, buccal tablets, troches, capsules, elixirs,  
20 suspensions, syrups, wafers, and the like. Such compositions and preparations should contain at least 0.1% of active compound. The percentage of the compositions and preparations may, of course, be varied and may conveniently be between about 2 to about 6% of the weight of the unit. The amount of active compound in such therapeutically useful compositions is such that a suitable  
25 dosage will be obtained. Preferred compositions or preparations according to the present invention are prepared so that an oral dosage unit form contains between about 50 and 300 mg of active compound.

          The tablets, troches, pills, capsules and the like may also contain the  
30 following: A binder such as gum tragacanth, acacia, corn starch or gelatin; excipients such as dicalcium phosphate; a disintegrating agent such as corn starch, potato starch, alginic acid and the like; a lubricant such as magnesium stearate; and a sweetening agent such as sucrose, lactose or saccharin may be added or a flavoring agent such as peppermint, oil of wintergreen, or cherry  
35 flavoring. When the dosage unit form is a capsule, it may contain, in addition to materials of the above type, a liquid carrier. Various other materials may be present as coatings or to otherwise modify the physical form of the dosage unit.

For instance, tablets, pills, or capsules may be coated with shellac, sugar or both. A syrup or elixir may contain the active compound, sucrose as a sweetening agent, methyl and propylparabens as preservatives, a dye and flavoring such as cherry or orange flavor. Of course, any material used in  
5 preparing any dosage unit form should be pharmaceutically pure and substantially non-toxic in the amounts employed. In addition, the active compound may be incorporated into sustained-release preparations and formulations.

10 The active compound may also be administered parenterally or intraperitoneally. Solutions of the active compound as a free base or pharmacologically acceptable salt can be prepared in water suitably mixed with a surfactant such as hydroxypropylcellulose. Dispersion can also be prepared in glycerol, liquid polyethylene glycols, and mixtures thereof and in oils. Under  
15 ordinary conditions of storage and use, these preparations contain a preservative to prevent the growth of microorganisms.

The pharmaceutical forms suitable for injectable use include sterile aqueous solutions or dispersions and sterile powders for the extemporaneous  
20 preparation of sterile injectable solutions or dispersions. In all cases the form must be sterile and must be fluid to the extent that easy syringability exists. It may be stable under the conditions of manufacture and storage and must be preserved against the contaminating action of microorganisms such as bacteria and fungi. The carrier can be a solvent or dispersion medium containing, for  
25 example, water, ethanol, polyol (for example, glycerol, propylene glycol, and liquid polyethylene glycol, and the like), suitable mixtures thereof, and vegetable oils. The proper fluidity can be maintained, for example, by the use of a coating such as lecithin, by the maintenance of the required particle size in the case of dispersion and by the use of surfactants. The prevention of the action of  
30 microorganisms can be brought about by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, sorbic acid, thimerosal, and the like. In many cases, it will be preferable to include isotonic agents, for example, sugars or sodium chloride. Prolonged absorption of the injectable compositions of agents delaying absorption, for example, aluminum  
35 monostearate and gelatin.

Sterile injectable solutions are prepared by incorporating the active compound in the required amount in the appropriate solvent with various of the other ingredients enumerated above, as required, followed by filtered sterilization. Generally, dispersions are prepared by incorporating the various  
5 sterilized active ingredient into a sterile vehicle which contains the basic dispersion medium and the required other ingredients from those enumerated above. In the case of sterile powders for the preparation of sterile injectable solutions, the preferred methods of preparation are vacuum drying and the freeze drying technique which yield a powder of the active ingredient plus any  
10 additional desired ingredient from previously sterile-filtered solution thereof.

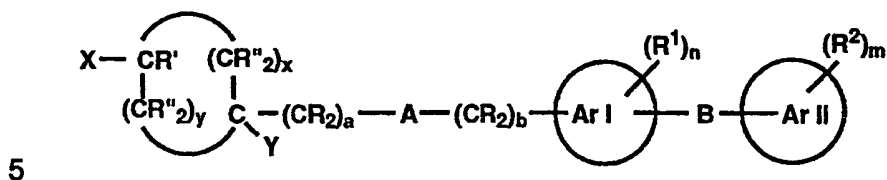
The therapeutic compounds of this invention may be administered to a mammal alone or in combination with pharmaceutically acceptable carriers, as noted above, the proportion of which is determined by the solubility and  
15 chemical nature of the compound, chosen route of administration and standard pharmaceutical practice.

The physician will determine the dosage of the present therapeutic agents which will be most suitable for prophylaxis or treatment and it will vary  
20 with the form of administration and the particular compound chosen, and also, it will vary with the particular patient under treatment. He will generally wish to initiate treatment with small dosages by small increments until the optimum effect under the circumstances is reached. The therapeutic dosage will generally be from about 0.1 to about 100 mg/kg/dy, and preferably from about  
25 10 mg to about 1000 mg day, or from about 0.1 mg to about 50 mg/kg of body weight per day and preferably from about 0.1 to about 20 mg/kg of body weight per day and may be administered in several different dosage units. Higher dosages on the order of about 2x to about 4x are required for oral  
administration.

30

## THE CLAIMS:

1. A compound of the formula:



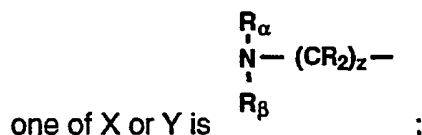
where:

Ar I and Ar II are independently a mono- or di- aryl or heteroaryl ring;

- 10 A is O, S, NR, SO, SO<sub>2</sub>, NR-C=O, O=C-NR, RC=CR, C/C or a bond;

- B is (CR<sub>2</sub>)<sub>e</sub>, O, S, NR, SO, SO<sub>2</sub>, NR-C=O, O=C-NR, RC=CR, C/C, O=C or a bond when Ar II is mono- aryl or heteroaryl and (CR<sub>2</sub>)<sub>e</sub>, O, S, O=C, NR, SO, SO<sub>2</sub> or a bond when Ar II is di- aryl or heteroaryl;
- 15

one of X or Y is R<sup>m</sup>;



20

a and b are independently 0-4 and a + b = 0-4;

e is 1-2;

25

m and n are independently 0-2;

x is 1-6;

y is 0-2;

30

x + y is 1-6;



z is 0-3;

$R_{\alpha}$  is hydrogen or alkyl;

5  $R_{\beta}$  is hydrogen, alkyl or hydroxy;

$R'$  is hydrogen, alkyl, hydroxy, alkoxy or halo;

10  $R''$  is hydrogen, alkyl, hydroxy, hydroxyalkyl, alkoxy, alkoxyalkyl or halo;

$R'''$  is hydrogen, alkyl, hydroxy or carboxy;

one set of vicinal  $R''$  and  $R'''$  groups may form a double bond  
when  $x+y=3-6$ ;

15

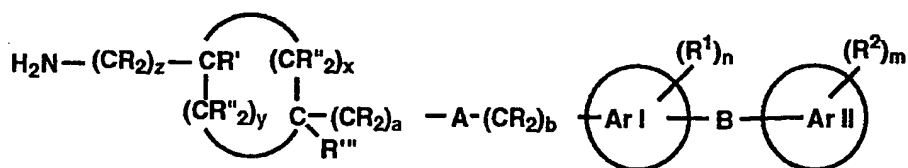
$R$  is hydrogen or alkyl;

$R^1$  and  $R^2$  are independently hydrogen, alkyl, alkoxy, hydroxy, halo,  
haloalkyl, nitro, amino or mono- or di-alkylamino; and

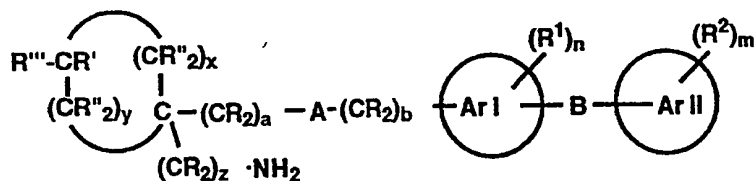
20

its stereoisomers, enantiomers, diastereoisomers and racemic mixtures; or  
pharmaceutically acceptable salts thereof.

25 2. A compound according to claim 1 of the formula:



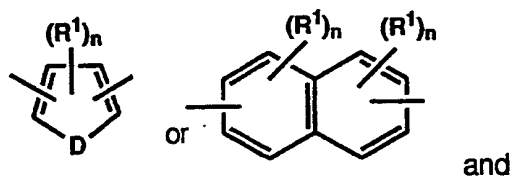
30 3. A compound according to Claim 1 of the formula:



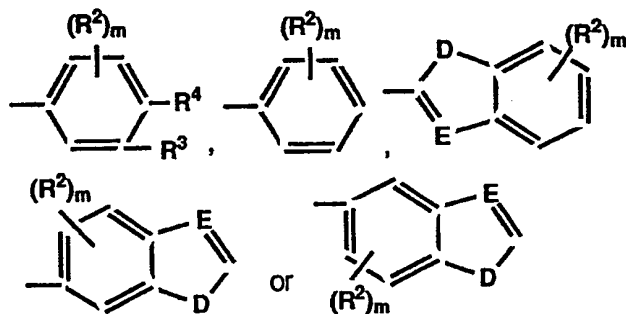
4. A compound according to claim 2 where:

5

Ar I is



Ar II is



10

where:

D is  $\text{CH}_2$ ,  $\text{CH}=\text{CH}$ , O, S or N-R;

15

E is CH, N or CH-O;

$R^1$  and  $R^2$  are independently hydrogen, alkyl, alkoxy, hydroxy, halo or trifluoromethyl; and

20

$R^3$  and  $R^4$  together are  $\text{O}-\text{CH}_2-\text{CH}_2$ ,  $\text{O}-\text{CH}_2-\text{CH}_2-\text{CH}_2$ ,  $\text{O}-\text{CH}_2-\text{O}$  or  $\text{O}-\text{CH}_2-\text{CH}_2-\text{O}$ ;

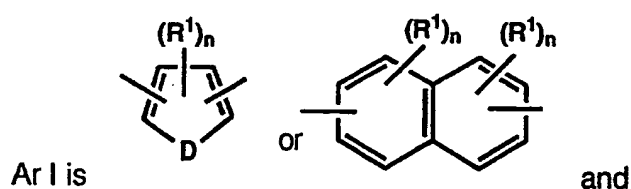
$R'$  is hydrogen, alkyl or hydroxy;

$R''$  is hydrogen, alkyl or hydroxy; and

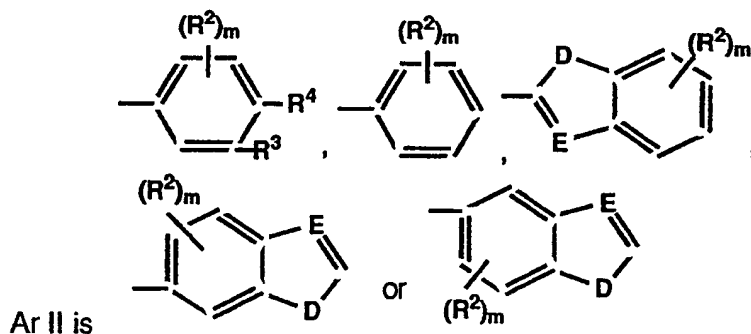
$R'''$  is hydrogen, alkyl, hydroxy or carboxy;

5

5. A compound according to claim 3 where:



10



where:

D is  $\text{CH}_2$ ,  $\text{CH}=\text{CH}$ , O, S or N-R;

15

E is CH, N or CH-O;

$R^1$  and  $R^2$  are independently hydrogen, alkyl, alkoxy, hydroxy, halo or trifluoromethyl; and

20

$R^3$  and  $R^4$  together are  $\text{O}-\text{CH}_2-\text{CH}_2$ ,  $\text{O}-\text{CH}_2-\text{CH}_2-\text{CH}_2$ ,  $\text{O}-\text{CH}_2-\text{O}$  or  $\text{O}-\text{CH}_2-\text{CH}_2-\text{O}$ ;

$R'$  is hydrogen, alkyl or hydroxy;

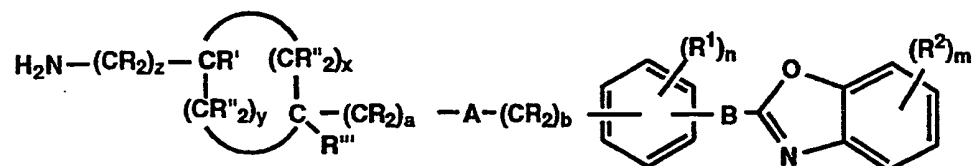
25

$R''$  is hydrogen, alkyl or hydroxy; and

$R'''$  is hydrogen, alkyl, hydroxy or carboxy;

6. A compound according to Claim 4 of the formula

5



where:

A is O, S, NR, CH=CH or a bond;

10

B is  $(CR_2)_e$ , O=C, O, S, NR or a bond;

a and b are independently 0-3 and  $a + b = 0-3$ ;

e is 1-2;

15

m and n are independently 0-2;

x is 1-6;

20

y is 0-2;

$x + y$  is 1-6;

z is 0-3;

25

$R'$  is hydrogen, alkyl or hydroxy;

$R''$  is hydrogen, alkyl or hydroxy;

30

$R'''$  is hydrogen, alkyl, hydroxy or carboxy;

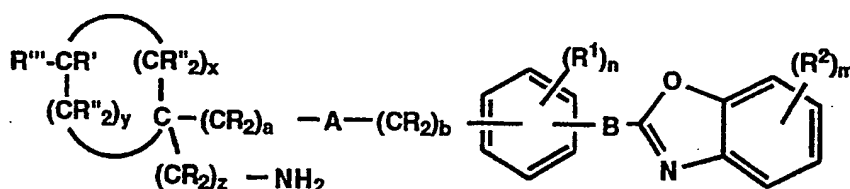
one set of vicinal  $R''$  and  $R'''$  groups may form a double bond  
when  $x+y=3-6$ ;

R is hydrogen or alkyl; and

R<sup>1</sup> and R<sup>2</sup> are independently hydrogen, alkyl, alkoxy, hydroxy, halo or trifluoromethyl.

5

7. A compound according to Claim 5 of the formula



10 where:

A is CH=CH, O, S, NR or a bond;

B is (CR<sub>2</sub>)<sub>e</sub>, O=C, O, S, NR or a bond;

15 a and b are independently 0-3 and a + b = 0-3;

e is 1-2;

m and n are independently 0-2;

20

x is 1-6;

y is 0-2;

25

x + y is 1-6;

z is 0-3;

R' is hydrogen, alkyl or hydroxy;

30

R<sup>a</sup> is hydrogen, alkyl or hydroxy;

R''' is hydrogen, alkyl, hydroxy or carboxy;

one set of vicinal R" and R''' groups may form a double bond  
when  $x+y=3-6$ ;

R is hydrogen or alkyl; and

5

R<sup>1</sup> and R<sup>2</sup> are independently hydrogen, alkyl, alkoxy, hydroxy, halo or  
trifluoromethyl.

10 8. A compound according to Claim 6 where:

A is O or a bond;

B is a bond;

15

a is 0-1;

b is 0-1;

20 m and n are independently 0-1;

x is 1-6;

y is 0-2;

25

$x + y$  is 1-6;

z is 0-1;

30 R, R', R'', R''' and R<sup>1</sup> are hydrogen or hydroxy; and

R<sup>2</sup> is hydrogen, alkyl, alkoxy or halo.

35 9. A compound according to Claim 7 where:

A is O or a bond;

B is a bond;

a is 0-1;

5

b is 0-1;

m and n are independently 0-1;

10

x is 1-6;

y is 0-2;

x + y is 1-6;

15

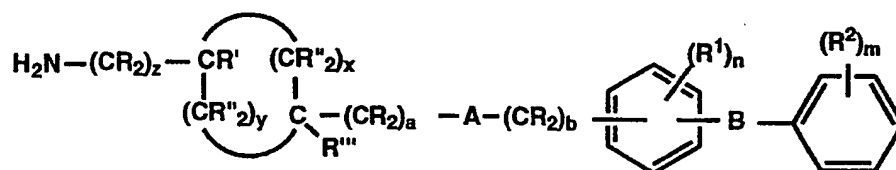
z is 0-1;

R, R', R'', R''' and R<sup>1</sup> are hydrogen or hydroxy; and

20

R<sup>2</sup> is hydrogen, alkyl, alkoxy or halo.

10. A compound according to Claim 4 of the formula



25

where:

A is O, S, NR, CH=CH or a bond;

B is (CR<sub>2</sub>)<sub>e</sub>, O, S, NR, O=C, CH=CH or a bond;

30

a and b are independently 0-3 and a + b = 0-3;

e is 1-2;

m and n are independently 0-2;

x is 1-6;

5 y is 0-2;

x + y is 1-6;

z is 0-3;

10

R' is hydrogen, alkyl or hydroxy;

R" is hydrogen, alkyl or hydroxy;

15

R''' is hydrogen, alkyl, hydroxy or carboxy;

one set of vicinal R" and R''' groups may form a double bond  
when x+y=3-6;

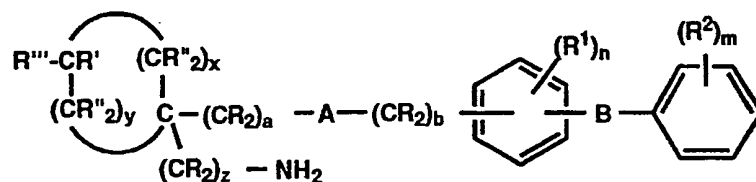
20

R is hydrogen or alkyl; and

R<sup>1</sup> and R<sup>2</sup> are independently hydrogen, alkyl, alkoxy, hydroxy, halo or  
trifluoromethyl.

25

11. A compound according to Claim 5 of the formula



where:

30

A is CH=CH, O, S, NR or a bond;

B is (CR2)e, O, S, NR, O=C, CH=CH or a bond;

a and b are independently 0-3 and a + b = 0-3;



e is 1-2;

m and n are independently 0-2;

5

x is 1-6;

y is 0-2;

10

x + y is 1-6;

z is 0-3;

15

R' is hydrogen, alkyl or hydroxy;

R'' is hydrogen, alkyl or hydroxy;

R''' is hydrogen, alkyl, hydroxy or carboxy;

20

one set of vicinal R'' and R''' groups may form a double bond  
when x+y=3-6;

R is hydrogen or alkyl; and

25

R<sup>1</sup> and R<sup>2</sup> are independently hydrogen, alkyl, alkoxy, hydroxy, halo or  
trifluoromethyl.

30

12. A compound according to Claim 10 where:

A is O;

B is CH=CH or a bond;

35

a is 0-1;

b is 0-1;

m and n are independently 0-1;

x is 1-6;

5

y is 0-2;

x + y is 1-6;

10

z is 0-1;

R, R', R'', R''' and R<sup>1</sup> are hydrogen or hydroxy; and

R<sup>2</sup> is hydrogen, alkyl, alkoxy or halo.

15

13. A compound according to Claim 11 where:

A is O;

20

B is CH=CH or a bond;

a is 0-1;

25

b is 0-1;

m and n are independently 0-1;

x is 1-6;

30

y is 0-2;

x + y is 1-6;

z is 0-1;

35

R, R', R'', R''' and R<sup>1</sup> are hydrogen or hydroxy; and

R<sup>2</sup> is hydrogen, alkyl, alkoxy or halo.

14. A compound according to Claim 6 which is trans-2-[4-(benzoxazol-2-yl)benzyloxy]cyclohexylamine or a pharmaceutically acceptable salt thereof.
15. A compound according to Claim 6 which is cis-2-[(4-benzoxazol-2-yl)benzyloxy]cyclohexylamine or a pharmaceutically acceptable salt thereof.
16. A compound according to Claim 6 which is trans-2-amino-[4-(benzoxazol-2-yl)benzyloxy]cyclopentane or a pharmaceutically acceptable salt thereof.
17. A compound according to Claim 6 which is trans-2-(4-[benzoxazol-2-yl]benzyloxy)cycloheptylamine or a pharmaceutically acceptable salt thereof.
18. A compound according to Claim 7 which is 2 $\alpha$ -hydroxy-3 $\beta$ [(4-benzoxazol-2-yl)benzyloxy]cyclohexyl-1 $\beta$ -amine or a pharmaceutically acceptable salt thereof.
19. A compound according to Claim 7 which is 2 $\beta$ -hydroxy-3 $\beta$ [(4-benzoxazol-2-yl)benzyloxy]cyclohexyl-1 $\alpha$ -amine or a pharmaceutically acceptable salt thereof.
20. A compound according to Claim 7 which is 1-amino-[4-(benzoxazol-2-yl)benzyloxymethyl]cyclopentane or a pharmaceutically acceptable salt thereof.
21. A method of lowering or maintaining reduced cholesterol levels in a patient requiring such treatment which comprises administering to such patient a squalene synthase inhibitor effective amount of a compound of the formula according to Claim 1.

22. A method for inhibiting cholesterol biosynthesis which comprises administering to a patient in need of such inhibition a squalene synthase  
5 inhibiting effective amount of a compound according to Claim 1.
23. A method according to Claim 22 where the patient is in need of a  
10 hypocholesterolemic or hypolipidemic agent.
24. A method according to Claim 23 for treating atherosclerosis.
- 15 25. A pharmaceutical composition comprising a squalene synthetase inhibitor effective amount of a compound according to Claim 1 in admixture with a pharmaceutical carrier.
- 20 26. A pharmaceutical composition according to Claim 25 which further includes an HMG CoA reductase inhibitor.

1/1

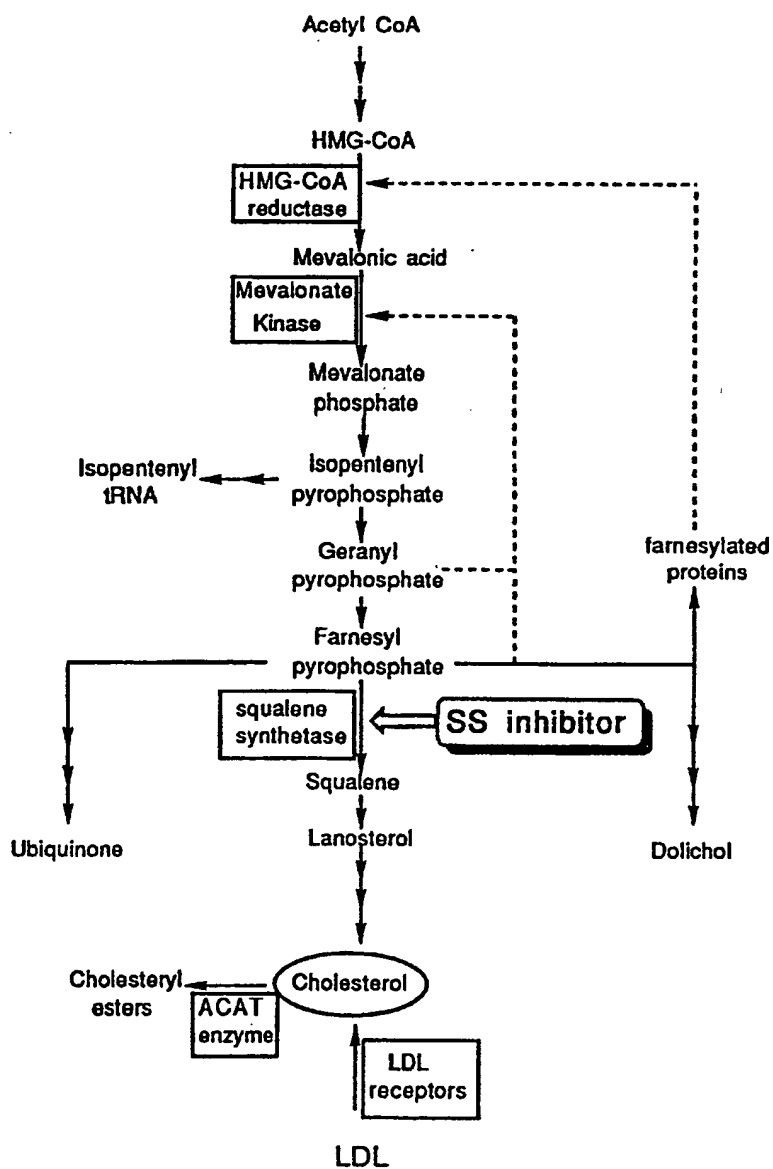


Figure 1. Cholesterol Biosynthetic Pathway

# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US93/12638

## A. CLASSIFICATION OF SUBJECT MATTER

IPC(5) :A61K 31/42; C07D 263/56

US CL :548/217, 219; 514/375

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/375; 548/217, 219

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CAS Online Structure Search

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US, A, 5,091,500 (LYSENKO et al.) 25 February 1992, see entire document.	1-26
Y	US, A, 4,826,838 (RICHARDSON et al.) 02 May 1989. See column line 65 to column 2 line 65 and claim 1.	1-5, 21-6
A	US, A, 5,135,935 (ALBERTS et al.) 04 August 1992.	1-26
A	US, A, 4,447,611 (KLAUVBERT et al.) 08 May 1984.	1-26
A	US, A, 4,681,883 (BROWN et al.) 21 July 1987.	1-26

☒ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	*T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
*A* document defining the general state of the art which is not considered to be part of particular relevance	*X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
*E* earlier document published on or after the international filing date	*Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
*L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*G* document member of the same patent family
*O* document referring to an oral disclosure, use, exhibition or other means	
*P* document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

26 APRIL 1994

Date of mailing of the international search report

MAY 12 1994

Name and mailing address of the ISA/US  
Commissioner of Patents and Trademarks  
Box PCT  
Washington, D.C. 20231

Facsimile No. (703) 305-3230

Authorized officer

D.G. DAUS ach

Telephone No. (703) 308-1235

## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US93/12638

## C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO, A, 92/12144 (SHIOTA et al.) 23 July 1992. See page 5 & 6 definitions for I when n is zero, compound 469, page 54.	1-5, 21-6
Y,P	WO, A, 93/09115 (HARRISON et al.) 13 May 1993. See entire document.	1-26
A	Jour. American Chem. Soc., Vol. 104, No. 25 issued 1982, SANDIFER et al. "SQUALENE SYNTHETASE: Inhibition by an Ammonium Analog of a Carbocationic Intermediate in the Conversion of Presqualene Pyrophosphate to Squalene", see pages 7376-8.	1-26
A	JOUR. AMER. CHEM. SOC., VOL. 111, NO. 10, issued 1989, POULTER et al. "SQUALENE SYNTHETASE. INHIBITION By AMMONIUM ANALOGUES OF CARBOCATIONIC INTERMEDIATES IN THE CONVERSION OF PRESQUALENE O. PHOSPHATE TO SQUALENE," pages 3734-39.	1-26
A	JOUR. MED. CHEM., VOL. 31, NO. 10, issued OCTOBER 1988, BILLER et al., "ISOPRENOID (PHOSPHINYLMETHYL) PHOSPHONATES AS INHIBITORS OF SQUALENE SYNTHETASE", pages 3734-39.	1-26
A	Jour. Med. Chem., Vol. 31, No. 10, issued October 1988, Biller et al. "ISOPRENOID (PHOSPHINYLMETHYL) PHOSPHONATES AS INHIBITORS OF SQUALENE SYNTHETASE", pages 1869-71.	1-26
A	Jour. of Pharmaceutical Sci., Vol. 80, No. 8, issued August 1991, STERLING et al. "QUATERNARY AND TERTIARY QUINUCLIDINE DERIVATIVES AS INHIBITORS OF CHOLINE UPTAKE", pages 785-9.	1-26
Y	WO, A, 92/15579 (NEUENSCHWANDER et al.) 17 September 1992. See claim 1 pages 54-5, where R1 and R3 may be (CH <sub>2</sub> ) <sub>n</sub> and B is CR'R'1	1-26

# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US93/12638

## Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:  
because they relate to subject matter not required to be searched by this Authority, namely:
  
2. ☒ Claims Nos.: 1-3, 21-26  
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:  
  
"Heteroaryl" cannot be completely searched in the PTO files, but these claims are searched to the extent they read on benzoxazoles.
  
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
  
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
  
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
  
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.  
☐ No protest accompanied the payment of additional search fees.